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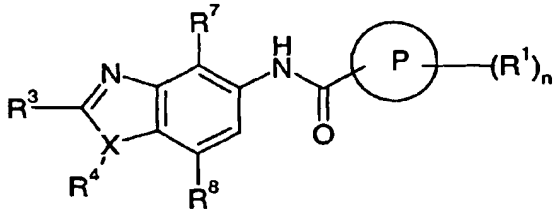
(54) Title: NEW HETEROCYCLIC AMIDES EXHIBITING AN INHIBITORY ACTIVITY AT THE VANILLOID RECEPTOR 1 (VR1).



(I)

(57) Abstract: The present invention relates to new compounds of formula I, wherein R^1 to R^8 are as defined as in formula I, or salts, solvates or solvated salts thereof, processes for their preparation and to new intermediates used in the preparation thereof, pharmaceutical compositions containing said compounds and to the use of said compounds in therapy.

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NEW HETEROCYCLIC AMIDES

FIELD OF THE INVENTION

The present invention relates to new compounds, to pharmaceutical compositions containing said compounds and to the use of said compounds in therapy. The present invention further relates to processes for the preparation of said compounds and to new intermediates used in the preparation thereof.

BACKGROUND OF THE INVENTION

Pain sensation in mammals is due to the activation of the peripheral terminals of a specialized population of sensory neurons known as nociceptors. Capsaicin, the active ingredient in hot peppers, produces sustained activation of nociceptors and also produces a dose-dependent pain sensation in humans. Cloning of the vanilloid receptor 1 (VR1 or TRPV1) demonstrated that VR1 is the molecular target for capsaicin and its analogues. (Caterina,M.J., Schumacher,M.A., et.al. Nature (1997) v.389 p 816-824). Functional studies using VR1 indicate that it is also activated by noxious heat , tissue acidification) and other inflammatory mediators (Tominaga,M., Caterina,M.J. et.al. Neuron (1998) v.21, p.531-543). Expression of VR1 is also regulated after peripheral nerve damage of the type that leads to neuropathic pain. These properties of VR1 make it a highly relevant target for pain and for diseases involving inflammation. While agonists of the VR1 receptor can act as analgesics through nociceptor destruction, the use of agonists, such as capsaicin and its analogues, is limited due to their pungency, neurotoxicity and induction of hypothermia. Instead, agents that block the activity of VR1 should prove more useful. Antagonists would maintain the analgesic properties, but avoid pungency and neurotoxicity side effects. Compounds with VR1 inhibitor activity are believed to be of potential use for the treatment and/or prophylaxis of disorders such as pain, especially that of inflammatory or traumatic origin such as arthritis, ischaemia, cancer, fibromyalgia, low back pain and post-operative pain (Walker et al J Pharmacol Exp Ther. (2003) Jan;304(1):56-62). In addition to this visceral pains such as chronic pelvic pain, cystitis, irritable bowel syndrome (IBS),

pancreatitis and the like, as well as neuropathic pain such as sciatia, diabetic neuropathy, HIV neuropathy, multiple sclerosis, and the like (Walker et al *ibid*, Rashid et al J Pharmacol Exp Ther. (2003) Mar;304(3):940-8), are potential pain states that could be treated with VR1 inhibitonThese compounds are also believed to be potentially useful for inflammatory disorders like asthma, cough, inflammatory bowel disease (IBD) (Hwang and Oh Curr Opin Pharmacol (2002) Jun;2(3):235-42). Compounds with VR1 blocker activity are also useful for itch and skin diseases like psoriasis and for gastro-esophageal reflux disease (GERD), emesis, cancer, urinary incontinence and hyperactive bladder (Yiangou et al BJU Int (2001) Jun;87(9):774-9, Szallasi Am J Clin Pathol (2002) 118: 110-21). VR1 inhibitors are also of potential use for the treatment and/or prophylaxis of the effects of exposure to VR1 activators like capsaicin or tear gas, acids or heat (Szallasi *ibid*).

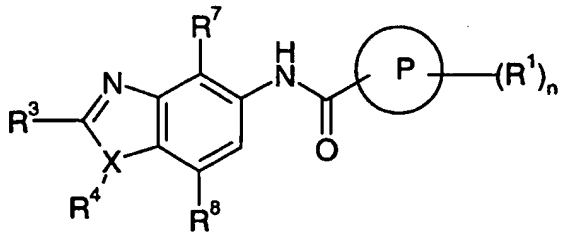
A further portential use relates to the treatment of tolerance to VR1 activators.

VR1 inhibitors may also be useful in the treatment of interstitial cystitis and pain related to interstitial cystitis.

DETAILED DESCRIPTION OF THE INVENTION

The object of the present invention is to provide compounds exhibiting an inhibitory activity at the vanilloid receptor 1 (VR1).

The present invention provides a compound of formula I



(I)

wherein:

ring P is C_{6-10}aryl, C_{3-7}cycloalkyl, C_{5-6}heteroaryl, which ring P may be fused with phenyl, C_{5-6}heteroaryl, C_{3-7}cycloalkyl or C_{3-7}heterocycloalkyl;

R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, $C_{1-6}alkyl$, $C_{2-6}alkenyl$, $C_{2-6}alkynyl$, $C_{1-6}haloalkyl$, $C_{1-6}haloalkylO$, $phenylC_{0-6}alkyl$, $C_{5-6}heteroarylC_{0-6}alkyl$, $C_{3-7}cycloalkylC_{0-6}alkyl$, $C_{3-7}heterocycloalkylC_{0-6}alkyl$, $C_{1-6}alkylOC_{0-6}alkyl$, $C_{1-6}alkylSC_{0-6}alkyl$ or $C_{1-6}alkylNC_{0-6}alkyl$;

n is 1, 2, 3, 4 or 5;

X is O or S, when

R^3 is H, $C_{1-6}alkyl$, $C_{1-6}haloalkyl$, $R^5OC_{1-6}alkyl$, R^5OCO, R^5CO, NR^5R^6CO, $NR^5R^6C_{0-6}alkyl$, $C_{2-6}alkenylOC_{0-6}alkyl$ or $hydroxyC_{1-6}alkyl$; and

R^4 is nil; or

X is N, when

R^3 is H, $C_{1-6}alkyl$, $C_{1-6}iodoalkyl$, $C_{1-6}bromoalkyl$, $C_{1-6}chloroalkyl$, $C_{1-6}alkylOC_{0-6}alkyl$, $R^5OC_{1-6}alkyl$, R^5CO, R^5CO2, NR^5R^6CO, $NR^5R^6C_{0-6}alkyl$ or $C_{2-6}alkenylOC_{0-6}alkyl$; and

R^4 is H, $C_{1-4}alkyl$, $hydroxyC_{1-6}alkyl$ or $C_{1-6}alkylOC_{1-6}alkyl$; or

X is N, when R^3 is $C_{1-6}fluoroalkyl$ or $hydroxyC_{1-2}alkyl$ and R^4 is H;

R^5 and R^6 are independently selected from H, $C_{1-6}alkyl$, $C_{6-10}aryl$, $C_{5-6}heteroaryl$, $C_{1-4}alkylSO_2$ and C_{1-3} alkylCO;

R^7 and R^8 are independently selected from H, $C_{1-6}alkyl$, halo, cyano, $C_{1-6}alkylOC_{0-6}alkyl$, OH, NO_2 and COR^9, $N(R^9)_2$;

R^9 is H or $C_{1-6}alkyl$;

and wherein any alkyl, alkylOalkyl, haloalkyl, haloalkylO, phenyl, heteroaryl, cycloalkyl or heterocycloalkyl group may be substituted with one or more A; and

A is OH, NO_2, $C_{1-6}alkylCO$, $C_{1-6}alkylO(CO)$, $N(R^9)_2$, R^9S, R^9SO_2, halo or $C_{1-6}alkylOC_{0-6}alkyl$,

or salts, solvates or solvated salts thereof.

One embodiment of the invention relates to the compound of formula I wherein

ring P is $C_{6-10}aryl$, $C_{5-6}heteroaryl$, which ring P may be fused with $C_{3-7}heterocycloalkyl$;

R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, $C_{1-6}alkyl$, $C_{2-6}alkenyl$, $C_{1-6}haloalkyl$, $C_{1-6}haloalkylO$, $phenylC_{0-6}alkyl$, $C_{3-7}heterocycloalkylC_{0-6}alkyl$, $C_{1-6}alkylOC_{0-6}alkyl$ or $C_{1-6}alkylSC_{0-6}alkyl$;

n is 1, 2 or 3;

X is O or S, when

R^3 is C_{1-6}alkyl, NR^5R^6CO, $NR^5R^6C_{0-6}$alkyl, C_{2-6}alkenylOC_{0-6}alkyl or hydroxyC_{1-6}alkyl; and

R^4 is nil; or

X is N, when

R^3 is H or C_{1-6}alkyl; and

R^4 is C_{1-4}alkyl or hydroxyC_{1-6}alkyl; or

X is N, when R^3 is C_{1-6}fluoroalkyl and R^4 is H;

R^5 and R^6 are independently selected from H, C_{6-10}aryl, C_{5-6}heteroaryl, C_{1-4}alkylSO_2 and C_{1-3} alkylCO;

R^7 and R^8 are independently selected from H, halo and cyano;

and wherein any alkyl, phenyl, heteroaryl group may be substituted with one or more A; and

A is OH, NO_2, halo or C_{1-6}alkylOC_{0-6}alkyl;

or salts, solvates or solvated salts thereof.

In one embodiment of the invention X is S and R^3 is C_{1-6}alkyl, NR^5R^6CO, $NR^5R^6C_{0-6}$alkyl, C_{2-6}alkenylOC_{0-6}alkyl or hydroxyC_{1-6}alkyl.

In another embodiment X is S and R^3 is methyl.

In a further embodiment X is S and R^3 is hydroxymethyl.

In one embodiment of the invention X is O and R^3 is C_{1-6}alkyl or hydroxyC_{1-6}alkyl.

In another embodiment X is O and R^3 is methyl.

In a further embodiment X is O and R^3 is hydroxymethyl.

In one embodiment of the invention X is N and R^3 is C_{1-6}alkyl and R^4 is C_{1-6}alkyl or hydroxyC_{1-6}alkyl.

In another embodiment R^3 is methyl and R^4 is methyl or 2-hydroxyethyl.

In a further embodiment X is N and R^3 is trifluoromethyl and R^4 is H

R^5 and R^6 may optionally be substituted by A. In one embodiment R^5 and R^6 are selected independently from the group consisting of H, methylsulfonyl, acetyl and substituted or unsubstituted heteroaryl such as pyrazole or pyridine.

One embodiment of the invention relates to the compound of formula I wherein R^3 is hydroxymethyl, allyloxymethyl, ethoxymethyl, methoxypyridinylaminomethyl, pyrazolylaminomethyl, aminomethyl, methylsulfonylaminomethyl, acetylaminomethyl, carboxamide, methyl, hydroxyethyl, nitrophenylaminomethyl, hydroxycarbonyl or methoxycarbonyl.

R^4 may be selected from the group consisting of H, C_{0-4}alkyl or hydroxyC_{1-6}alkyl.

In one embodiment of the invention P is substituted with 0, 1, 2, 3 or 4 groups R^1, wherein the number of R^1 substituents on the P ring is designated by the term n. In another embodiment of the invention n is 1 or 2.

Another embodiment of the invention relates to the compound of formula I wherein ring P is phenyl.
In a further embodiment ring P is phenyl and R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{1-6}haloalkyl, C_{1-6}haloalkylO, phenylC_{0-6}alkyl, C_{5-6}heteroarylC_{0-6}alkyl, C_{3-7}cycloalkylC_{0-6}alkyl, C_{3-7}heterocycloalkylC_{0-6}alkyl, C_{1-6}alkylOC_{0-6}alkyl, C_{1-6}alkylSC_{0-6}alkyl or C_{1-6}alkylNC_{0-6}alkyl optionally substituted with one or more A.

In yet another embodiment ring P is pyrazolyl, pyridine, benzdioxolane, furan, thiophene or naphthalene and R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{1-6}haloalkyl, C_{1-6}haloalkylO, phenylC_{0-6}alkyl, C_{5-6}heteroarylC_{0-6}alkyl, C_{3-7}cycloalkylC_{0-6}alkyl, C_{3-7}heterocycloalkylC_{0-6}alkyl, C_{1-6}alkylOC_{0-6}alkyl, C_{1-6}alkylSC_{0-6}alkyl or C_{1-6}alkylNC_{0-6}alkyl optionally substituted with one or more A.

Ring P may be substituted by R^1 on a nitrogen or carbon atom in ring P. Further, one atom on ring P may be substituted by two substituents R^1.

Any alkyl, alkylOalkyl, haloalkyl, haloalkylO, phenyl, heteroaryl, cycloalkyl or heterocycloalkyl group present in the substituents of the compounds of formula I may be substituted with one or more A. One embodiment of the invention relates to compounds of formula I wherein A is selected from the group consisting of OH, NO_2, halo or C_{1-6}alkylOC_{0-6}alkyl.

Another embodiment of the invention relates to compounds selected from the group consisting of

3-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,

2-tert-Butyl-5-methyl-2*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,

2-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide,

4-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide,

3,4-Dimethyl-*N*-(2-methyl-benzothiazol-5-yl)-benzamide,

2,2-Difluoro-benzo[1,3]dioxole-5-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-6-trifluoromethyl-nicotinamide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-4-propyl-benzamide,

3-Iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

2,5-Dimethyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

5-tert-Butyl-2-methyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

4-Bromo-3-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

3,4-Difluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

3-Chloro-2-fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

Pyridine-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2-Benzyl-5-tert-butyl-2*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

3-Fluoro-4-trifluoromethyl-*N*-(2-trifluoromethyl-1*H*-benzimidazol-5-yl)-benzamide,

2-Fluoro-5-trifluoromethyl-*N*-(2-trifluoromethyl-1*H*-benzimidazol-5-yl)-benzamide,

4-Chloro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

1-Phenyl-5-trifluoromethyl-1*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

1-Phenyl-5-propyl-1*H*-pyrazole-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2,3-Difluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,

3-Fluoro-4-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-methyl-benzooxazol-5-yl)-benzamide,

Biphenyl-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

3-Bromo-thiophene-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

4-Bromo-2-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

2-Chloro-3,4-dimethoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Amino-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-nitro-benzamide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-4-vinyl-benzamide,

4-Ethoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethylsulfanyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Dimethylamino-naphthalene-1-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2-Fluoro-6-iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethoxymethyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-4-trifluoromethoxy-benzamide, and

4-Chloro-3-fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

or salts, solvates or solvated salts thereof.

A further embodiment of the invention relates to compounds selected from the group consisting of

4-*tert*-Butyl-*N*-(2-formyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-hydroxymethyl-1,3-benzothiazol-5-yl)-benzamide,

5-(4-tert-butylbenzoylamino)-1,3-benzothiazol-2-ylcarboxylic acid, and

4-tert-Butyl-*N*-(2-methoxycarbonyl-1,3-benzothiazol-5-yl)-benzamide,

or salts, solvates or solvated salts thereof.

Yet another embodiment of the invention relates to compounds selected from the group consisting of

4-tert-Butoxy-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

4-Bromo-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-iodobenzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-morpholin-4-ylbenzamide,

N-{2-[(Allyloxy)methyl]-1,3-benzothiazol-5-yl}-4-morpholin-4-ylbenzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-propyl-1H-pyrazole-4-carboxamide,

1-tert-Butyl-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-3-methyl-1H-pyrazole-5-carboxamide,

4-(Ethoxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-1H-pyrazole-5-carboxamide,

4-Bromo-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-2-methylbenzamide,

4-tert-Butoxy-N-(2-methyl-1,3-benzoxazol-5-yl) benzamide,

N-(4-Bromo-2-methyl-1,3-benzothiazol-5-yl)-4-tert-butylbenzamide,

4-tert-Butyl-N-(4,7-dibromo-2-methyl-1,3-benzothiazol-5-yl)benzamide,

N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-(trifluoromethyl)-1H-pyrazole-4-carboxamide,

4-Iodo-N-(2-methyl-5-benzothiazolyl)benzamide,

4-(tert-Butoxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-(1,2-Dimethyl-1H-benzimidazol-5-yl)-4-iodobenzamide,

4-tert-Butyl-N-(2-{[(2-methoxypyridin-3-yl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide,

4-tert-Butyl-N-[2-(1-hydroxyethyl)-1,3-benzothiazol-5-yl]benzamide,

4-tert-Butyl-N-{2-[(1H-pyrazol-3-ylamino)methyl]-1,3-benzothiazol-5-yl}benzamide,

4-(1,1-Dimethylethyl)-N-[2-[[(4-nitrophenyl)amino]methyl]-5-benzothiazolyl]-benzamide,

N-[2-(Aminomethyl)-1,3-benzothiazol-5-yl]-4-tert-butylbenzamide,

4-tert-Butyl-N-(2-{[(methylsulfonyl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide,

N-{2-[(Acetylamino)methyl]-1,3-benzothiazol-5-yl}-4-tert-butylbenzamide,

5-[(4-tert-Butylbenzoyl)amino]-1,3-benzothiazole-2-carboxamide,

N-1,3-Benzothiazol-5-yl-4-tert-butylbenzamide,
4-Chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-propyl-1H-pyrazole-4-carboxamide,
1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-(trifluoromethyl)-1H-pyrazole-4-carboxamide,
N-(2,4-dimethyl-1,3-benzothiazol-5-yl)-4-(1-hydroxy-1-methylethyl)benzamide,
4-(Hydroxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamid,e and
4-tert-butyl-N-(4-cyano-2-methyl-1,3-benzothiazol-5-yl)benzamide,
or salts, solvates or solvated salts thereof.

One embodiment of the invention relates to compounds selected from the group consisting of.
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-[2,2,2-trifluoro-1-hydroxy-1-(trifluoromethyl)ethyl]benzamide,
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-isopropoxybenzamide,
4-Bromo-2-chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
4-Bromo-2-fluoro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(morpholin-4-ylmethyl)benzamide,
3-Fluoro-*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(trifluoromethyl)benzamide,
4-tert-butoxy-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
4-(tert-Butoxymethyl)-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide, and
4-*tert*-butyl-*N*-[2-(hydroxymethyl)-1,3-benzoxazol-5-yl]benzamide,
or salts, solvates or solvated salts thereof.

Listed below are definitions of various terms used in the specification and claims to describe the present invention.

For the avoidance of doubt it is to be understood that where in this specification a group is qualified by 'hereinbefore defined', 'defined hereinbefore' or 'defined above' the said

group encompasses the first occurring and broadest definition as well as each and all of the other definitions for that group.

For the avoidance of doubt it is to be understood that in this specification 'C_{1-6}' means a carbon group having 1, 2, 3, 4, 5 or 6 carbon atoms.

In this specification, unless stated otherwise, the term "alkyl" includes both straight and branched chain alkyl groups and may be, but are not limited to methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, s-butyl, t-butyl, n-pentyl, i-pentyl, t-pentyl, neo-pentyl, n-hexyl or i-hexyl, t-hexyl. The term C_{1-3} alkyl having 1 to 3 carbon atoms and may be methyl, ethyl, n-propyl, i-propyl or *tert*-butyl.

The term 'C_0' means a bond or does not excist. For example when R^4 is C_0alkyl, R^4 does not excist and "arylC_0alkyl" is equivalent with "aryl", "C_2aklylOC_0alkyl" is equivalent with "C_2alkylO".

In this specification, unless stated otherwise, the term "alkenyl" includes both straight and branched chain alkenyl groups. The term "C_{2-6}alkenyl" having 2 to 6 carbon atoms and one or two double bonds, may be, but is not limited to vinyl, allyl, propenyl, butenyl, crotyl, pentenyl, or hexenyl, and a butenyl group may for example be buten-2-yl, buten-3-yl or buten-4-yl.

In this specification, unless stated otherwise, the term "alkynyl" includes both straight and branched chain alkynyl groups. The term "C_{2-6}alkynyl" having 2 to 6 carbon atoms and one or two trippel bonds, may be, but is not limited to etynyl, propargyl, pentynyl or hexynyl and a butynyl group may for example be butyn-3-yl or butyn-4-yl.

In this specification, unless stated otherwise, the term "cycloalkyl" refers to an optionally substituted, saturated cyclic hydrocarbon ring system. The term "C_{3-7}cycloalkyl" may be cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl or cycloheptyl.

The term "heterocycloalkyl" denotes a 3- to 7-membered, non-aromatic, partially or completely saturated hydrocarbon group, which contains one ring and at least one heteroatom. Examples of said heterocycle include, but are not limited to pyridyl, pyrrolyl, furyl, thienyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, pyrazolyl, benzofuryl, indolyl, isoindolyl, benzimidazolyl, pyridazinyl, pyrimidinyl, pyrazinyl, tetrazolyl, triazolyl, pyrrolidinyl, pyrrolidonyl, piperidinyl, piperazinyl, morpholinyl, oxazolyl, 2-oxazolidonyl or tetrahydrofuranyl.

In this specification, unless stated otherwise, the term "aryl" refer to an optionally substituted monocyclic or bicyclic hydrocarbon unsaturated aromatic ring system. Examples of "aryl" may be, but are not limited to phenyl and naphthyl.

In this specification, unless stated otherwise, the term "heteroaryl" refer to an optionally substituted monocyclic or bicyclic unsaturated aromatic ring system containing at least one heteroatom selected independently form N, O or S. Examples of "heteroaryl" may be, but are not limited to pyridyl, pyrrolyl, furyl, thienyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, pyrazolyl, benzofuryl, indolyl, isoindolyl, benzimidazolyl, pyridazinyl, pyrimidinyl, pyrazinyl, tetrazolyl, triazolyl and oxazolyl.

In this specification, unless stated otherwise, the term "arylalkyl" and "heteroarylalkyl" refer to a substituent that is attached via the alkyl group to an aryl or heteroaryl group.

In this specification, unless stated otherwise, the term "halo" and "halogen" may be fluoro, iodo, chloro or bromo.

In this specification, unless stated otherwise, the term "haloalkyl" means an alkyl group as defined above, which is substituted with halo as defined above. The term "C_{1-6}haloalkyl" may include, but is not limited to fluoromethyl, difluoromethyl, trifluoromethyl, fluoroethyl, difluoroethyl or bromopropyl. The term "C_{1-6}haloalkylO" may include, but is not limited to fluoromethoxy, difluoromethoxy, trifluoromethoxy, fluoroethoxy or difluoroethoxy.

The present invention relates to the compounds of formula I as hereinbefore defined as well as to the salts, solvates or solvated salts thereof. Salts for use in pharmaceutical compositions will be pharmaceutically acceptable salts, but other salts may be useful in the production of the compounds of formula I.

A suitable pharmaceutically acceptable salt of the compounds of the invention is, for example, an acid-addition salt, for example an inorganic or organic acid. In addition, a suitable pharmaceutically acceptable salt of the compounds of the invention is an alkali metal salt, an alkaline earth metal salt or a salt with an organic base.

Other pharmaceutically acceptable salts and methods of preparing these salts may be found in, for example, Remington's Pharmaceutical Sciences (18th Edition, Mack Publishing Co.).

Some compounds of formula I may have chiral centres and/or geometric isomeric centres (E- and Z- isomers), and it is to be understood that the invention encompasses all such optical, diastereoisomeric and geometric isomers.

The invention also relates to any and all tautomeric forms of the compounds of formula I.

Methods of Preparation

Another aspect of the present invention provides processes for preparing compounds of formula I, or salts, solvates or solvated salts thereof.

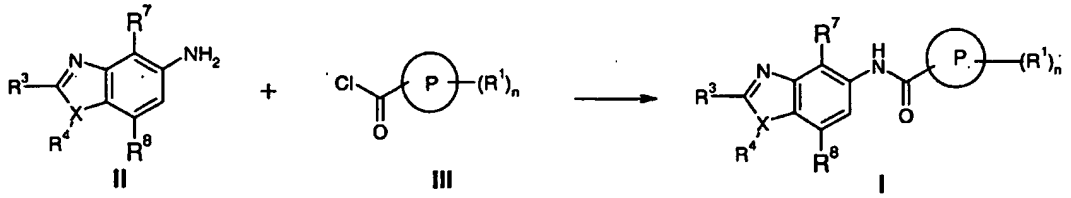
Throughout the following description of such processes it is to be understood that, where appropriate, suitable protecting groups will be added to, and subsequently removed from, the various reactants and intermediates in a manner that will be readily understood by one skilled in the art of organic synthesis. Conventional procedures for using such protecting groups as well as examples of suitable protecting groups are described, for example, in "Protective Groups in Organic Synthesis", T.W. Green, P.G.M. Wuts, Wiley-Interscience, New York, (1999). References and descriptions of other suitable reactions are described in textbooks of organic chemistry, for example, "Advanced Organic Chemistry", March, 4th ed. McGraw Hill (1992) or, "Organic Synthesis", Smith, McGraw Hill, (1994). For representative examples of heterocyclic chemistry see for example "Heterocyclic

Chemistry", J. A. Joule, K. Mills, G. F. Smith, 3rd ed. Chapman and Hall (1995), p. 189-224 and "Heterocyclic Chemistry", T. L. Gilchrist, 2nd ed. Longman Scientific and Technical (1992), p. 248-282.

The term "room temperature" and "ambient temperature" shall mean, unless otherwise specified, a temperature between 16 and 25 °C.

One embodiment of the invention relates to processes for the preparation of the compound of formula I, wherein R^1 to R^8, unless otherwise specified, are defined as in formula I, comprising;

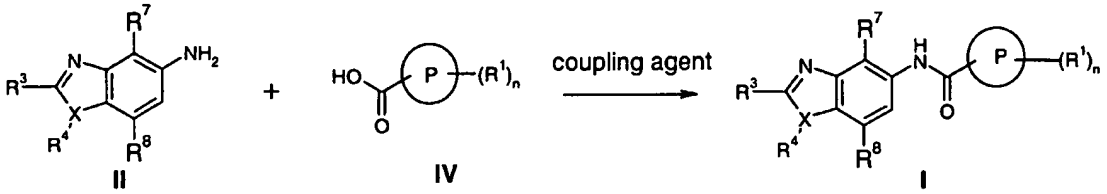
a) reaction of an aromatic amine of formula (II) with a properly substituted acyl chloride (III) optionally in the presence of a base:



II III I

This reaction may be performed in any manner known to the skilled person in the art. Suitable solvents to be used for this reaction may be halogenated hydrocarbons such as chloroform, dichloromethane and dichloroethane or aromatic and heteroaromatic compounds such as benzene, toluene, xylene, pyridine and lutidine or ethers such as ethyl ether, tetrahydrofuran and dioxan or any mixtures thereof. Catalysts such as heteroaromatic bases like pyridine and lutidine or tertiary amines like triethylamine, *N*-methylmorpholine and ethyl diisopropylamine or polymer bound tertiary amines like *N*,*N*-(diisopropyl)aminomethylpolystyrene resin may be used as well. The temperature may be between -40 and 40°C and the reaction time may be between 0.5 and 30 h.

b) reaction of an aromatic amine of formula (II) with a properly substituted acid (IV) in the presence of a coupling agent (activator) like for example 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide hydrochloride.



II IV I

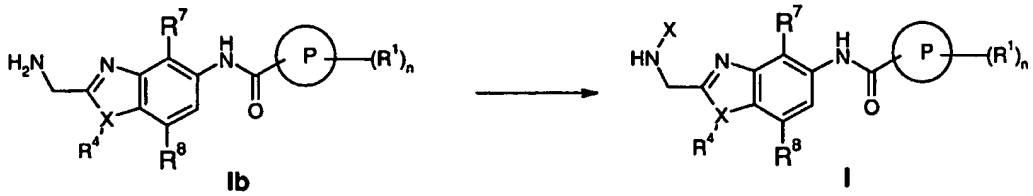
Suitable solvents to be used for this reaction may be tertiary amides such as dimethylformamide and dimethylacetamide, halogenated hydrocarbons such as chloroform, dichloromethane and dichloroethane or aromatic and heteroaromatic compounds such as benzene, toluene, xylene, pyridine and lutidine or ethers such as ethyl ether, tetrahydrofuran and dioxan or any mixtures thereof. Catalysts such as heteroaromatic bases like pyridine and lutidine or tertiary amines like triethylamine, *N*-methylmorpholine and ethyl diisopropylamine may be used as well. The temperature may be between 10 and 60°C and the reaction time may be between 3 and 30 h.

c) reaction of an hydroxymethyl derivative Ia with methanesulfonyl chloride followed by treatment with ammonia.

Ia Ib

The mesylation step is carried out using halogenated hydrocarbons such as chloroform, dichloromethane and dichloroethane as a solvent and a tertiary amine like triethylamine, *N*-methylmorpholine and ethyl diisopropylamine as a base in a temperature range between –20 and 30 °C. The amination step step is carried out using a solution of ammonia in an alcohol like ethanol or in an aprotic solvent like dioxane or in water.

d) reaction of an aminomethyl derivative Ib with an acyl chloride or a sulfonyl chloride

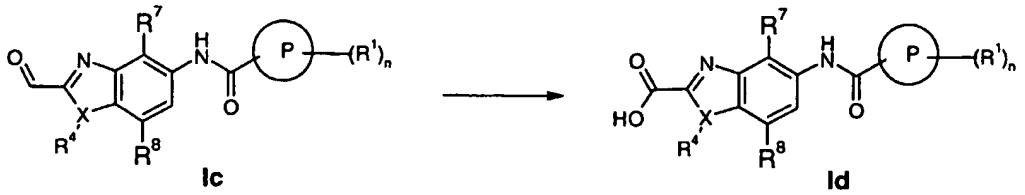


Ib I

X = acyl, sulfonyl

The reaction conditions are similar to the ones described for the mesylation step in part c).

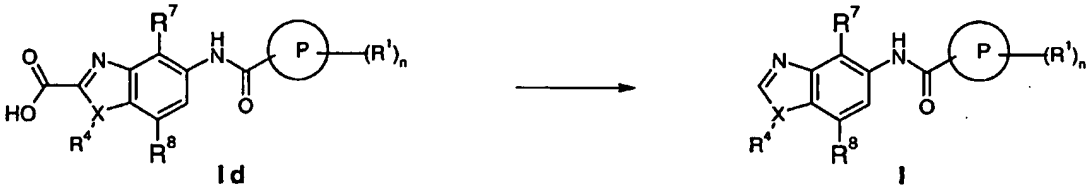
e) oxidation of the aldehyde Ic to the corresponding carbonic acid Id



Ic Id

For the oxidation purpose a mixture of sodium chlorite and sulfamic acid in water may be employed

f) decarboxylation of the carbonic acid Id

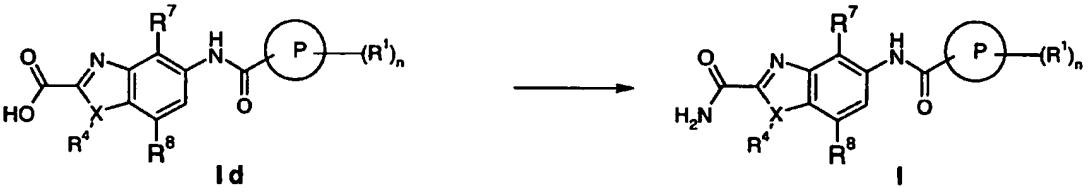


Id I

g) esterification of the carbonic acid Id

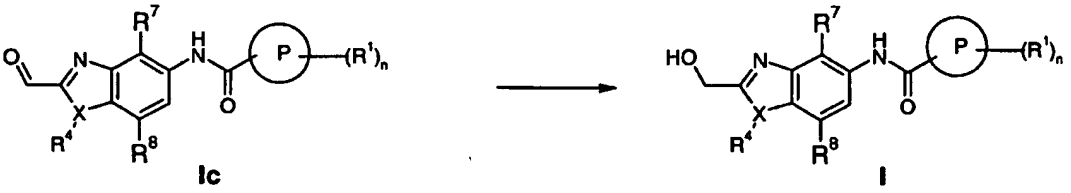
Id

h) amidation of the carbonic acid Id



Id I

i) reduction of the aldehyde Ic to a corresponding primary alcohol



Ic I

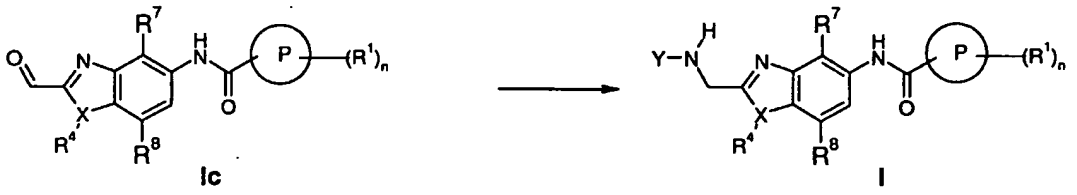
As a suitable reductive agent sodium borohydride may be used in a solvent like methanol or another alcohol or its mixture with water in a temperature range between -10 and 40°C

j) treatment of the aldehyde Ic with organometallic reagent leading to secondary alcohols

Ic I

Organometalic reagent may be a magnesium derivatives like methylmagnesium bromide or organolithium compound like methyllithium and a suitable solvent may be chosen from a range of aprotic inert solvents like diethyl ether, tetrahydrofuran, benzene, etc.

k) reductive amination of the aldehyde Ic

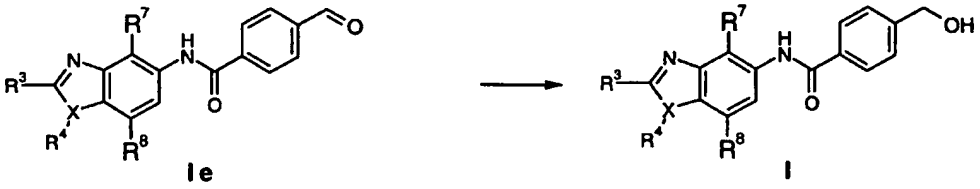


Ic I

Y is aryl, heteroaryl

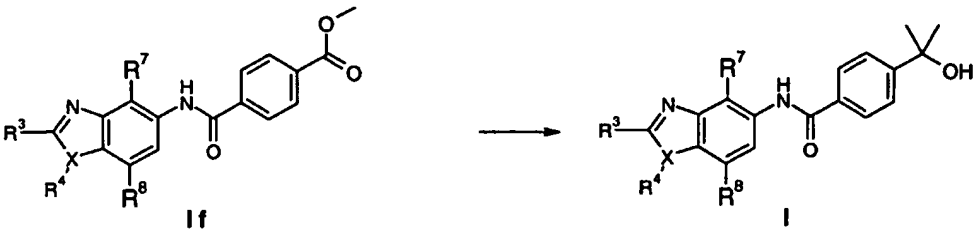
in process i) any primary amine may be used together with an appropriate reductive agent for example decaborane or sodium cyanoborohydride. Both protic and aprotic solvents, for example, alcohols, water, terahydrofuran and mixtures thereof are suitable and the temperature range is between 0 and 40°C.

l) reduction of the aldehyde Ie to a corresponding primary alcohol as in part i)



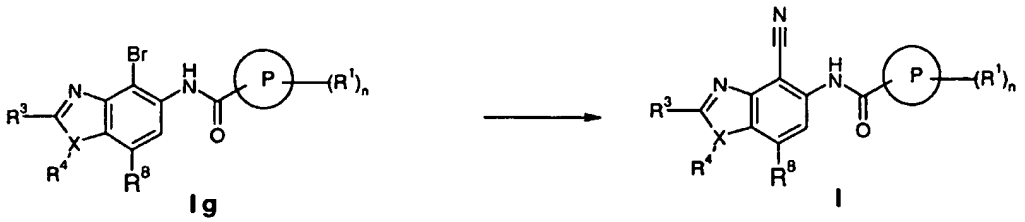
Ie I

m) treatment of the methyl ester If with organometallic reagent leading to tertiary alcohols in a similar way to the process described in part j)



If I

n) reaction of the bromo derivative Ig with a cyanation reagent



Ig I

As a cyanation reagent copper (I) cyanide may be used in an aprotic polar solvent having high boiling point, like dimethyl formamide, at elevated temperature in a range between 150 and 270°C

o) oxidation of the 2-methyl derivative Ih and subsequent reduction of the intermediary aldehyde to the 2-hydroxymethyl derivative Ii

Ih Ii

The oxidation step is accomplished by using an appropriate oxidative reagent for example, magnesium dioxide, chromium trioxide or selenium dioxide. Suitable solvents to be used for this reaction may be ketones such as acetone and butan-2-one, or halogenated hydrocarbons such as chloroform, dichloromethane and dichloroethane or any mixtures thereof. The temperature may be between 0 and 80°C and the reaction time may be between 3 and 50 h. The subsequent reduction is typically carried out using sodium borohydride in methanol.

Abbreviations

alloc	allyloxycarbonyl
DCE	dichloroethane
DCM	dichloromethane
DMAP	dimethylaminopyridine
EDC	1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
HATU	O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate
HPLC	high performance liquid chromatography
LC	liquid chromatography

MsCl	methanesulfonyl chloride
MS	mass spectometry
ret. time	retention time
TFA	trifluroacetic acid

A further embodiment of the invention relates to compounds
allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate,
4-tert-Butyl-N-(2-formyl-1,3-benzothiazol-5-yl)-benzamide, and
4-Bromo-2-methyl-benzothiazol-5-ylamine, and
4-chloro-2-methyl-benzothiazole-5-ylamine.

Another embodiment relates to the used of these compounds as intermediates in the preparation of the compound of formula I.

Pharmaceutical composition

According to one embodiment of the present invention there is provided a pharmaceutical composition comprising as active ingredient a therapeutically effective amount of the compound of formula I, or salts, solvates or solvated salts thereof, in association with one or more pharmaceutically acceptable diluents, excipients and/or inert carriers.

The composition may be in a form suitable for oral administration, for example as a tablet, pill, syrup, powder, granule or capsule, for parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion) as a sterile solution, suspension or emulsion, for topical administration e.g. as an ointment, patch or cream or for rectal administration e.g. as a suppository.
In general the above compositions may be prepared in a conventional manner using one or more conventional excipients, pharmaceutical acceptable diluents and/or inert carriers.
Suitable daily doses of the compounds of formula I in the treatment of a mammal, including man, are approximately 0.01 to 250 mg/kg bodyweight at peroral administration and about 0.001 to 250 mg/kg bodyweight at parenteral administration.
The typical daily dose of the active ingredient varies within a wide range and will depend

on various factors such as the relevant indication, severity of the illness being treated, the route of administration, the age, weight and sex of the patient and the particular compound being used, and may be determined by a physician.

Examples of pharmaceutical composition

The following illustrate representative pharmaceutical dosage forms containing a compound of formula I, or salts, solvates or solvated salts thereof, (hereafter compound X), for preventive or therapeutic use in mammals:

(a): Tablet	**mg/tablet**
Compound X	100
Lactose	182.75
Croscarmellose sodium	12.0
Maize starch paste (5% w/v paste)	2.25
Magnesium stearate	3.0

(b): Capsule	**mg/capsule**
Compound X	10
Lactose	488.5
Magnesium stearate	1.5

(c): Injection	**(50 mg/ml)**
Compound X	5.0% w/v
1M Sodium hydroxide solution	15.0% v/v
0.1M Hydrochloric acid	(to adjust pH to 7.6)
Polyethylene glycol 400	4.5% w/v
Water for injection	up to 100%

The above compositions may be obtained by conventional procedures well known in the pharmaceutical art.

Medical use

Surprisingly, it has been found that the compounds according to the present invention are useful in therapy. The compounds of formula I, or salts, solvates or solvated salts thereof, as well as their corresponding active metabolites, exhibit a high degree of potency and selectivity for individual vanilloid receptor 1 (VR1) groups. Accordingly, the compounds of the present invention are expected to be useful in the treatment of conditions associated with excitatory activation of vanilloid receptor 1 (VR1).

The compounds may be used to produce an inhibitory effect of VR1 in mammals, including man.

VR1 are highly expressed the peripheral nervous system and in other tissues. Thus, it is expected that the compounds of the invention are well suited for the treatment of VR1 mediated disorders.

The compounds of formula I are expected to be suitable for the treatment of acute and chronic pain, acute and chronic neuropathic pain and acute and chronic inflammatory pain.

Examples of such disorder may be selected from the group comprising arthritis, fibromyalgia, low back pain, post-operative pain, visceral pains like chronic pelvic pain, cystitis, including interstitial cystitis, bowel syndrome (IBS), pancreatitis, ischeamic, sciatia, diabetic neuropathy, multiple sclerosis, HIV neuropathy, asthma, cough and inflammatory bowel disease (IBD).

Further relevant disorders may be selected from the group comprising gastro-esophageal reflux disease (GERD), psoriasis, cancer, emesis, urinary incontinence and hyperactive bladder.

Other relevant disorders are related to respiratory diseases and may be selected from the group comprising asthma, chronic obstructive lung disease and emphysema, lung fibrosis and interstitial lung disease.

The VR1 inhibitor(s) may be administrated by either an oral or inhaled route. The respiratory disease may be an acute and chronic illness and may be related to infection(s) and/or exposure to environmental pollution and/or irritants.

The compounds of formula I may also be used as antitoxin to treat (over-) exposure to VR1 activators like capsaicin, tear gas, acids or heat. Regarding heat, there is a potential use for VR1 antagonists in (sun-) burn induced pain, or inflammatory pain resulting from brun injuries.

The compounds may further be used for treatment of tolerance to VR1 activators.

One embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, in therapy.

Another embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of VR1 mediated disorders.

A further embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of acute and chronic pain disorders.

Yet another embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of acute and chronic neuropathic pain.

Yet a further embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of acute and chronic inflammatory pain.

One embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of arthritis, fibromyalgia, low back pain, post-operative pain, visceral pains like chronic pelvic pain, cystitis, IBS, pancreatitis or ischeamic.

Another embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of sciatia, diabetic neuropathy, multiple sclerosis or HIV neuropathy.

A further embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of asthma, cough, IBD, psoriasis, gastro-esophageal reflux disease (GERD), psoriasis, cancer, emesis, urinary incontinence or hyperactive bladder.

Yet another embodiment of the invention relates to the use of the compounds of formula I as hereinbefore defined, for treatment of interstitial cystitis and pain related to interstitial cystitis.

Yet a further embodiment of the invention relates to the use of the compound of formula I as hereinbefore defined, for the treatment of respiratory diseases selected from the group comprising asthma, chronic obstructive lung disease and emphysema, lung fibrosis and interstitial lung disease.

One embodiment of the invention relates to the use of the compound of formula I as hereinbefore defined, in the manufacture of a medicament for treatment of VR1 mediated disorders and for treatment of acute and chronic pain disorders, acute and chronic neuropathic pain and acute and chronic inflammatory pain and any other disorder mentioned above.

Another embodiment of the invention relates to a method of treatment of VR1 mediated disorders and acute and chronic pain disorders, acute and chronic neuropathic pain and acute and chronic inflammatory pain, and respiratory diseases, and any other disorder mentioned above, comprising administrering to a mammal, including man in need of such treatment, a therapeutically effective amount of the compounds of formula I, as hereinbefore defined.

A further embodiment of the invention relates to a pharmaceutical composition comprising a compound of formula I as hereinbefore defined, for use in treatment of VR1 mediated disorders and for treatment of acute and chronic pain disorders, acute and chronic neuropathic pain and acute and chronic inflammatory pain and any other disorder mentioned above.

In the context of the present specification, the term "therapy" and "treatment" includes prevention and prophylaxis, unless there are specific indications to the contrary. The terms "treat","therapeutic" and "therapeutically" should be construed accordingly.

In this specification, unless stated otherwise, the term "inhibitor" and "antagonist" mean a compound that by any means, partly or completely, blocks the transduction pathway leading to the production of a response by the ligand.

The term "disorder", unless stated otherwise, means any condition and disease associated with vanilloid receptor activity.

Non- Medical use

In addition to their use in therapeutic medicine, the compounds of formula I, or salts, solvates or solvated salts thereof, are also useful as pharmacological tools in the development and standardisation of *in vitro* and *in vivo* test systems for the evaluation of the effects of inhibitors of VR1 related activity in laboratory animals such as cats, dogs, rabbits, monkeys, rats and mice, as part of the search for new therapeutics agents.

Examples

The invention will now be illustrated by the following non-limiting examples.

General methods

All starting materials are commercially available or described in the literature. The ^1H NMR spectra were recorded on Brucker at 400 MHz. The mass spectra were recorded utilising electrospray (LC-MS; LC:Waters 2790, column XTerra MS C_8 2.5 µm 2.1X30 mm, buffer gradient H_2O+0.1%TFA:CH_3CN+0.04%TFA, MS: micromass ZMD// ammonium acetate buffer) ionisation techniques.

Synthesis of aromatic amines as starting materials employed in amide bond-forming reactions in examples 1-17.

Allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate

A. *tert*-Butyl (2-methyl-1,3-benzothiazol-5-yl)carbamate.

A mixture of Et_3N (100 mL), di-*tert*-butyl dicarbonate (58.3 g, 267 mmol) and 5-amino-2-methylbenzothiazole (22.0 g, 134 mmol) in MeOH (300 mL) was stirred at 65 °C for 2 hours and room temperature for 18 hours. The mixture was concentrated under reduced pressure, and the residue was diluted with DCM and washed with a 1M solution of HCl. The organic phase was dried with Na_2SO_4, filtered and evaporated under reduced pressure to yield the carbamate derivative. $R_f = 0.45$ (hexanes:EtOAc, 1:1); MS [M+] calc. 264.0 found 264.9.

B. *tert*-Butyl [2-(hydroxymethyl)-1,3-benzothiazol-5-yl]carbamate.

SeO_2 (45.0 g, 402 mmol) was ground to a fine powder and added to a solution of the carbamate in dioxane (300 mL). The mixture was kept under a N_2 atmosphere and heated to 70 °C for 18 hours with vigorous stirring. The mixture was quickly filtered and the solid was washed with hot dioxane. The filtrate was concentrated under reduced pressure to yield the aldehyde. $R_f = 0.56$ (hexanes:EtOAc, 1:1). The crude aldehyde was dissolved in MeOH (300 mL) and $NaBH_4$ (15.21 g, 402 mmol) was added portion-wise. The mixture was stirred for 2 hours and then diluted with 1M NaOH. The mixture was evaporated to dryness, dissolved in DCM, washed with a saturated solution of $NaHCO_3$, dried with Na_2SO_4, filtered and concentrated under reduced pressure. $R_f = 0.09$ (hexanes:EtOAc, 2:1); MS [M+] calc. 280.0 found 280.9.

C. Allyl {5-[(*tert*-butoxycarbonyl)amino]-1,3-benzothiazol-2-yl}methyl carbonate.

The primary alcohol was dissolved in DCM (300 mL), and allylchloroformate (16.2 mg, 134 mmol) was added followed by DMAP (14.2 g, 140 mmol). The mixture was stirred for 3 hours, and the solvent was evaporated. MS [M+] calc. 364.0 found 364.9.

D. Allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate.

The alloc-protected derivative was dissolved in DCM (300 mL), and TFA (100 mL) was added. The mixture was stirred for 18 hours, and then concentrated under reduced pressure. The product was purified by flash chromatography on silica gel eluting with mixtures of heptane and EtOAc (4:1, 7:3 and 1:1) to yield an off-white powder (6.6 g, 25 mmol). ^{1}H NMR (400 MHz, CHLOROFORM-D) δ ppm 4.71 (d, *J*=5.86 Hz, 2 H) 5.30 (dd, *J*=10.35, 1.17 Hz, 1 H) 5.37 (q, *J*=3.0, 1.50 Hz, 1 H) 5.42 (q, *J*=3.0, 1.50 Hz, 1 H) 5.51 (s, 2 H) 5.95 (m, 1 H) 7.10 (s, 1 H) 7.63 (s, 1 H) 7.70 (d, *J*=8.01 Hz, 1 H); ^{13}C NMR (101 MHz, DMSO-D6) δ ppm 65.9, 68.5, 110.7, 117.5, 118.7, 122.9, 126.9, 131.9, 140.5, 153.4, 153.9, 166.7; MS [M+] calc. 264.0 found 264.8.

Allyl (5-amino-4-chloro-1,3-benzothiazol-2-yl)methyl carbonate

Allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate (500 mg, 1.89 mmol) was dissolved in DCM (19.0 mL) and N-chlorosuccinimide (253 mg, 1.89 mmol) was added. The mixture was stirred at ambient temperature until the reaction appeared complete by LC-MS. The solution was concentrated under reduced pressure and purified by flash chromatography using mixtures of hexanes and EtOAc (4:1, 2:1) as an eluent to yield the title product (429 mg, 1.44 mmol, 76%). ^{1}H NMR (400 MHz, chloroform-D) δ ppm 2.77 (s, 2H) 4.71 (dt, *J*=5.86, 2.73, 1.37 Hz, 2 H) 5.27 - 5.46 (m, 2 H) 5.57 (s, 2 H) 5.89 - 6.05 (m, 1 H) 6.92 (d, *J*=8.59 Hz, 1 H) 7.55 (d, *J*=8.59 Hz, 1 H).

4-Bromo-2-methyl-1,3-benzothiazol-5-ylamine and 4,6-Dibromo-2-methyl-benzothiazol-5-ylamine

5-Amino-2-methylbenzothiazole (2.45 g, 14.9 mmol) and Br_2 (2.38 g, 14.9 mmol) were mixed in $CHCl_3$ (60.0 mL) and stirred for 45 minutes. 28% NH_4OH (20.0 mL) was added,

and the aqueous phase was extracted with DCM. The combined organic phases were dried with $MgSO_4$, filtered and evaporated. The products were separated from by flash chromatography on silica gel eluting with mixtures of hexanes and EtOAc (4:1) to yield 4-bromo-2-methyl-1,3-benzothiazol-5-ylamine: LC ret. time 1.13 minutes (Column: Phenomonex Polar, Gradient: 10-95% B, Flow rate:1.75 mL/min, Column temperature: 40 °C, Mobile phase: A - 0.1% TFA in H_2O, B - 0.1% TFA in MeCN), MS [M+] calcd. 242.0, found 242.0; and 4,6-dibromo-2-methyl-benzothiazol-5-ylamine: LC ret. time 1.64 minutes MS [M+] calcd. 322.0, found 322.0

5-Amino-1,3-benzothiazole-2-carbaldehyde

Manganese dioxide (10 mmol) was added to a solution of 5-amino-2-methylbenzothiazole (2 mmol) in acetone (20 mL). The mixture was refluxed for 24 h. After cooling to ambient temperature, the mixture was filtered and concentrated *in vacuo* to afford crude 5-amino-2-formylbenzothiazole as a yellow oil which was used for the next step without further purification. MS [M+] calc. 178.2, found 179

Example 1

4-tert-Butoxy-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide.

A. Synthesis of the O-alloc protected derivative

Allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate (see above) (97.0 mg, 0.370 mmol) and 4-*tert*-butoxybenzoic acid (71.0 mg, 0.370 mmol) were mixed in a mixture of DCM (5.00 mL) and DMF (5.00 mL) with EDC (220 mg, 1.15 mmol) and DMAP (236 mg, 1.15 mmol).). The mixture was stirred for 18 hours, and the solvents were evaporated. The residue was dissolved in DCM and washed with a saturated solution of $NaHCO_3$. The mixture was dried with Na_2SO_4, filtered and concentrated under reduced pressure. The crude product was purified by Gilson reverse phase HPLC (Luna 15 u, C18 (2) 250 mm X 21.2 mm), eluting with mixtures of H_20 and MeCN with 0.1% TFA the O-alloc protected derivative of the title compound: MS [M+] calc. 440.0 found 440.9.

B. Deprotection

The product obtained in Part A was treated with a solution of $Pd(OAc)_2$ (10.0 mg), PPh_3 (20.0 mg) and Et_3SiH (176 mg, 1.52 mmol, 0.240 mL) in a mixture of THF (4.00 mL) and DMF (4.00 mL). The mixture was stirred at room temperature until the reaction appeared complete by TLC analysis, and the solvents were evaporated. The crude product was purified by Gilson HPLC (Luna 15 u, C18 (2), 250 mm X 21.2 mm) eluting with mixture of MeCN and H_2O containing 1%TFA to yield the title product. ^1H NMR (400 MHz, methanol-D4) δ ppm 1.42 (s, 9 H) 4.95 (s, 2 H) 7.12 (m, 2 H) 7.71 (dd, J=8.79, 1.17 Hz, 1 H) 7.91 (m, 2 H) 7.96 (d, J=8.79 Hz, 1 H) 8.40 (s, 1 H); MS [M+H] calc. 357.1 found 357.0; Anal. found C 64.61% H 5.58% N 6.65%.

Example 2

4-Bromo-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide.

4-Bromobenzoylchloride (0.4 mmol) was dissolved in DCM and DMAP (0.4 mmol) was added. The mixture was stirred for 10 minutes and then allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate (100 mg, 0.38 mmol) was added. The mixture was stirred until the reaction appeared complete by TLC analysis and NaOH (1M) was added. The aqueous phase was extracted with DCM. The organic phases were collected, dried with Na_2SO_4, filtered and concentrated under reduced pressure. Purification by HPLC afforded the O-alloc protected derivative of the title compound: MS [M+1] calc. 448.0 found 448.4. Deprotection according to the procedure described in Example 1, part B afforded the title compound; δ ppm 4.79 (s, 2H) 7.55 (d, J=8.3 Hz, 3 H) 7.74 (d, J=8.4 Hz, 2 H) 7.80 (d, J=8.7 Hz, 1 H) 8.25 (s, 1 H); MS [M+H] calc. 363.2 found 363.0.

Compounds in the following examples were synthesized according to the amide bond-forming procedures described in the examples 1 or 2 starting from an appropriate aromatic amine, either commercially available or synthesized according to the procedures described above, and an appropriately substituted commercially available aromatic acid or an aromatic acyl chroride. Where appropriate the amide bond-forming procedures were followed by the deprotection as described in Example 1

Example number	Name	MW calcd	MW found [M+1]	^{1}H NMR
3	*N*-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-iodobenzamide	411.1	411.0	(600 MHz, methanol-D_4) δppm 4.79 (s, 2H) 7.55 (d, J=8.3 Hz, 3 H) 7.74 (d, J=8.4 Hz, 2 H) 7.80 (d, J=8.7 Hz, 1 H) 8.25 (s, 1 H)
4	N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-morpholin-4-ylbenzamide	370.1	370.0	(400 MHz, DMSO-D_6) δ ppm 3.24 (m, 4 H), 3.73 (m, 4 H) 4.83 (d, J=6.05 Hz, 2 H) 6.22 (t, J=5.96 Hz, 1 H) 7.02 (d, J=9.18 Hz, 2 H) 7.76 (dd, J=8.69, 2.05 Hz, 1 H) 7.90 (d, J=9.18 Hz, 2 H) 7.98 (d, J=8.79 Hz, 1 H) 8.42 (d, J=1.95 Hz, 1 H) 10.13 (s, 1 H)
5 (no deprotection step required)	N-{2-[(Allyloxy)methyl]-1,3-benzothiazol-5-yl}-4-morpholin-4-ylbenzamide	410.2	410.0	(400 MHz, methanol-D_4) δ ppm 3.09 (m, 4 H), 3.73 (m, 4 H) 4.58 (d, J=6.05 Hz, 2 H) 4.90 (s, 2 H) 5.16 (m, 2 H) 5.99 (m, 1 H) 6.70 (m, 2 H) 7.22 (m, 3 H) 7.62 (d, J=1.95 Hz, 1 H) 7.87 (d, J=8.59 Hz, 1 H)
6	N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-propyl-1H-pyrazole-4-carboxamide	393.1	392.9	(400 MHz, methanol-D_4) δ ppm 3.09 (m, 4 H), 3.73 (m, 4 H) 4.58 (d, J=6.05 Hz, 2 H) 4.90 (s, 2 H) 5.16 (m, 2 H) 5.99 (m, 1 H) 6.70 (m, 2 H) 7.22 (m, 3 H) 7.62 (d, J=1.95

				Hz, 1 H) 7.87 (d, J=8.59 Hz, 1 H)
7	1-tert-Butyl-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-3-methyl-1H-pyrazole-5-carboxamide	345.1	345.0	(400 MHz, methanol-D_4) δ ppm 1.71 (s, 9 H) 2.52 (s, 3 H) 4.95 (s, 2 H) 6.61 (s, 1 H) 7.69 (dd, J=8.69, 1.85 Hz, 1 H) 7.93 (d, J=8.79 Hz, 1 H) 8.43 (d, J=1.95 Hz, 1 H)
8	4-(Ethoxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	343.1	343.0	(400 MHz, methanol-D_4) δ ppm 1.24 (t, J=7.03 Hz, 3 H) 3.58 (q, J=7.03 Hz, 2 H) 4.58 (s, 2 H) 4.95 (s, 2 H) 7.48 (d, J=8.59 Hz, 2 H) 7.71 (dd, J=8.69, 2.05 Hz, 1 H) 7.92 (s, 1 H) 7.94 (d, J=7.81 Hz, 2 H) 8.41 (d, J=1.95 Hz, 1 H)
9	N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-1H-pyrazole-5-carboxamide	335.1	335.0	(400 MHz, chloroform-D) δ ppm 2.78 (s, 3 H) 6.79 (d, J=1.76 Hz, 1 H) 7.35 (m, 5 H) 7.60 (m, 3 H) 7.89 (s, 1 H)
10	4-Bromo-*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-2-methylbenzamide	377.0	377.0	(400 MHz, methanol-D_4) δ ppm 2.45 (s, 3 H) 4.95 (s, 2 H) 7.40 (d, J=8.20 Hz, 1 H) 7.45 (m, 1 H) 7.49 (s, 1 H) 7.66 (dd, J=8.69, 2.05 Hz, 1 H) 7.93 (d, J=8.59 Hz, 1 H) 8.40 (d, J=1.95 Hz, 1 H)
11	4-tert-Butoxy-N-(2-methyl-1,3-benzoxazol-5-yl) benzamide	325.1	325.2	(400 MHz, chloroform-D) δppm 1.41 (s, 9 H) 2.63 (s, 3 H) 7.07 (m, 2 H) 7.43 (d, J=8.79 Hz, 1 H) 7.58 (dt,

				J=8.79, 2.15 Hz, 1 H) 7.82 (m, 2 H) 7.89 (d, J=1.95 Hz, 1 H) 7.98 (s, 1 H)
12	N-(4-Bromo-2-methyl-1,3-benzothiazol-5-yl)-4-tert-butylbenzamide	403.0	403.0	(400 MHz, chloroform-D) δ ppm 1.38 (s, 9 H) 2.90 (s, 3 H) 7.56 (d, J=8.59 Hz, 2 H) 7.79 (d, J=8.79 Hz, 1 H) 7.93 (d, J=8.59 Hz, 2 H) 8.63 (d, J=8.98 Hz, 1 H)
13	4-*tert*-Butyl-*N*-(4,7-dibromo-2-methyl-1,3-benzothiazol-5-yl)benzamide	481.0	480.7	(400 MHz, methanol-D_4) δ ppm 1.37 (s, 9 H) 2.85 (s, 3 H) 7.58 (d, J=8.40 Hz, 2 H) 7.97 (d, J=8.40 Hz, 2 H) 8.31 (s, 1 H)
14	*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-(trifluoromethyl)-1*H*-pyrazole-4-carboxamide	419.1	419.0	(400 MHz, methanol-D4) δ ppm 4.95 (m, 2 H), 7.59 (m, 6 H), 7.96 (m, 1 H), 8.14 (m, 1 H), 8.37 (m, 1 H).
15	4-Iodo-*N*-(2-methyl-5-benzothiazolyl)benzamide	395.0	394.8	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H) 7.64 (d, *J*=8.59 Hz, 2 H) 7.73 (dd, *J*=8.59, 1.95 Hz, 1 H) 7.80 (m, 1 H) 7.86 (d, *J*=8.59 Hz, 2 H) 7.94 (m, 1 H) 8.14 (d, *J*=1.95 Hz, 1 H)
16	4-(*tert*-Butoxymethyl)-*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	371.1	371.0	(400 MHz, methanol-D_4) δ ppm 1.30 (s, 9 H) 4.54 (s, 2 H) 4.94 (s, 2 H) 7.48 (d, J=8.59 Hz, 2 H) 7.71 (dd, J=8.69, 2.05 Hz, 1 H) 7.92 (m, 3 H)

				8.39 (d, J=1.56 Hz, 1 H)
17	*N*-(1,2-Dimethyl-1*H*-benzimidazol-5-yl)-4-iodobenzamide	392.0	392.0	(400 MHz, DMSO-D_6) δ ppm 2.75 (s, 3 H) 3.87 (s, 3 H) 7.75 (d, J=8.59 Hz, 2 H) 7.80 (t, J=2.05 Hz, 1 H) 7.84 (d, J=8.98 Hz, 1 H) 7.92 (d, J=8.40 Hz, 2 H) 8.32 (t, J=2.15 Hz, 1 H) 10.58 (s, 1 H)
18	*N*-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-[2,2,2-trifluoro-1-hydroxy-1-(trifluoromethyl)ethyl]benzamide	450.1	451.0	(400 MHz, methanol-D4) δ ppm 4.95 (s, 2 H) 7.72 (d, J=8.59 Hz, 1 H) 7.90 (d, J=8.40 Hz, 2 H) 7.96 (d, J=8.59 Hz, 1 H) 8.04 (d, J=8.59 Hz, 2 H) 8.40 - 8.46 (m, 1 H) 10.43 (s, 1 H)
19	N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-isopropoxybenzamide	343.1	343.0	(400 MHz, methanol-D4) δ ppm 1.31 (d, J=6.05 Hz, 6 H) 4.65 (dt, J=11.96, 6.03 Hz, 1 H) 4.93 (s, 2 H) 6.95 (d, J=8.98 Hz, 2 H) 7.66 (dd, J=8.59, 1.76 Hz, 1 H) 7.85 - 7.91 (m, 2 H) 8.36 (s, 1 H)
20	4-Bromo-2-chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	397.0	396.7	(400 MHz, DMSO-D6) δ ppm 4.82 (s, 2 H) 6.23 (t, J=12.11, 6.05 Hz, 1 H) 7.58 (d, J=8.20 Hz, 1 H) 7.63 (dt, J=8.79, 3.32, 2.15 Hz, 1 H) 7.68 (dd, J=8.20, 1.95 Hz, 1 H) 7.88 (d, J=1.95 Hz, 1 H) 8.01 (d, J=8.59 Hz, 1 H) 8.35 (t, J=1.66 Hz, 1 H) 10.70 (s, 1 H)

21	4-Bromo-2-fluoro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	381.0	381.0	(400 MHz, chloroform-D) δ ppm 5.09 (s, 2 H) 7.43 (d, J=11.52 Hz, 1 H) 7.50 (dd, J=8.40, 1.56 Hz, 1 H) 7.71 (s, 1 H) 7.88 (d, J=8.59 Hz, 1 H) 8.11 (t, J=8.49 Hz, 1 H) 8.33 (s, 1 H) 8.52 (s, 1 H)
22	*N*-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(morpholin-4-ylmethyl)benzamide	384.1	384.0	(400 MHz, methanol-D4) δ ppm 3.18 - 3.46 (m, 4 H) 3.75 (s, 2 H) 3.94 - 4.17 (m, 2 H) 4.47 (s, 2 H) 4.96 (s, 2 H) 7.65 - 7.75 (m, 3 H) 7.98 (d, J=8.79 Hz, 1 H) 8.10 (d, J=8.20 Hz, 2 H) 8.42 (d, J=1.56 Hz, 1 H)
23	3-Fluoro-*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(trifluoromethyl)benzamide	371.0	371.0	(400 MHz, methanoL-D4) δ ppm 4.95 (s, 2 H) 7.73 (dd, J=8.69, 1.86 Hz, 1 H) 7.83 - 8.01 (m, 4 H) 8.43 (d, J=1.76 Hz, 1 H)
24	4-tert-butoxy-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	391.1	391.0	(400 MHz, methanol-D4) δ ppm 1.42 (s, 9 H) 4.98 (s, 2 H) 7.13 (d, J=8.59 Hz, 2 H) 7.75 (d, J=8.59 Hz, 1 H) 7.88 - 7.98 (m, 3 H)
25	4-(tert-Butoxymethyl)-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide	405.1	405.0	(400 MHz, methanol-D4) δ ppm 1.26 - 1.36 (m, 9 H) 4.57 (s, 2 H) 4.98 (s, 2 H) 5.48 (s, 1 H) 7.51 (d, J=8.20 Hz, 2 H) 7.75 (d, J=8.59 Hz, 1 H) 7.97 (dd, J=8.40, 2.73 Hz, 3 H)

26	3-Fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide	354.3	355.0	
27	2-tert-Butyl-5-methyl-2H-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	328.4	329	
28	2-Fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide	354.3	355.0	
29	2-Fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide	354.3	354.0	
30	4-Fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide	354.3	354.0	
31	3,4-Dimethyl-N-(2-methyl-benzothiazol-5-yl)-benzamide	296.4	297.1	
32	2,2-Difluoro-benzo[1,3]dioxole-5-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	348.3	349	
33	N-(2-Methyl-1,3-benzothiazol-5-yl)-6-trifluoromethyl-nicotinamide	337.3	338	

34	N-(2-Methyl-1,3-benzothiazol-5-yl)-4-propyl-benzamide	310.4	311.1	
35	3-Iodo-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	394.23	394.9	
36	2,5-Dimethyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	286.35	287	
37	5-tert-Butyl-2-methyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	328.43	329.1	
38	4-Bromo-3-methyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	361.26	360.99	
39	3,4-Difluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	304.32	305	
40	3-Chloro-2-fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	320.77	321	
41	Pyridine-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	269.33	270	
42	2-Benzyl-5-tert-butyl-2H-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	404.54	405.1	

43	3-Fluoro-4-trifluoromethyl-N-(2-trifluoromethyl-1H-benzimidazol-5-yl)-benzamide	391.25	392	
44	2-Fluoro-5-trifluoromethyl-N-(2-trifluoromethyl-1H-benzimidazol-5-yl)-benzamide	391.25	392	
45	4-Chloro-N-(2-methyl-benzothiazol-5-yl)-benzamide	302.8	302.9	(400 MHz, DMSO-D6) δ ppm 2.8 (s, 3 H) 7.6 (d, J=8.6 Hz, 2 H) 7.8 (d, J=9.1 Hz, 1 H) 8.0 (m, 3 H) 8.4 (s, 1 H) 10.5 (s, 1 H)
46	1-Phenyl-5-trifluoromethyl-1H-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	402.4	402.9	(400 MHz, DMSO-D6) δ ppm 2.8 (s, 3 H) 7.5 (m, 2 H) 7.6 (m, 3 H) 7.7 (d, J=9.1 Hz, 1 H) 8.0 (d, J=8.6 Hz, 1 H) 8.3 (m, 2 H) 10.7 (s, 1 H)
47	1-Phenyl-5-propyl-1H-pyrazole-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	376.5	376.9	(400 MHz, chloroform-D) δ ppm 0.8 (t, J=7.3 Hz, 3 H) 1.5 (m, 2 H) 2.8 (s, 3 H) 2.9 (m, 2 H) 7.4 (m, 2 H) 7.4 (m, 3 H) 7.7 (s, 1 H) 7.7 (m, 2 H) 7.9 (s, 1 H) 8.0 (m, 1 H)
48	2,3-Difluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide	372.3	372.7	(400 MHz, chloroform-D) δ ppm 2.8 (s, 3 H) 7.5 (t, J=7.3 Hz, 1 H) 7.7 (dd, J=8.6, 2.0 Hz, 1 H) 7.8 (d, J=8.6 Hz, 1 H) 8.0 (t, J=7.6 Hz, 1 H) 8.2

				(d, J=2.0 Hz, 1 H) 8.4 (d, broad, 1 H)
49	3-Fluoro-4-methyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	300.4	300.8	
50	4-tert-Butyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	324.5	325.2	(400 MHz, chloroform-D) δ ppm 1.4 (s, 9 H) 2.8 (s, 3 H) 7.5 (d, J=8.6 Hz, 2 H) 7.8 (m, 2 H) 7.8 (d, J=8.6 Hz, 2 H) 7.9 (s, 1 H) 8.1 (m, 1 H)
51	4-Ethyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	296.4	297.2	
52	4-tert-Butyl-N-(2-methyl-1,3-benzooxazol-5-yl)-benzamide	308.4	309	
53	Biphenyl-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	344.4	345	
54	3-Bromo-thiophene-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	353.3	354	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H) 7.11 (d, J=5.27 Hz, 1 H) 7.54 (d, J=5.27 Hz, 1 H) 7.73 (dd, J=8.79, 2.15 Hz, 1 H) 7.80 (d, J=8.59 Hz, 1 H) 8.20 (d, J=1.95 Hz, 1 H) 8.98 (s, 1 H)
55	4-Bromo-2-methyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	361.3	362	(400 MHz, chloroform-D) δ ppm 2.50 (s, 3 H) 2.84 (s, 3 H) 7.42 (d, J=19.14 Hz, 2 H) 7.64 (s, 1 H) 7.77 (m, 3 H) 8.10 (s, 1 H)

56	4-tert-Butoxy-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	340.4	341	(400 MHz, chloroform-D) δ ppm 1.41 (m, 9 H), 2.84 (s, 3 H), 7.09 (d, J=8.79 Hz, 2 H), 7.78 (m, 2 H), 7.83 (d, J=8.98 Hz, 2 H), 7.93 (s, 1 H), 8.12 (m, 1 H)
57	2-Chloro-3,4-dimethoxy-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	362.8	363	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H), 3.90 (s, 3 H), 3.94 (s, 3 H), 6.94 (d, J=8.79 Hz, 1 H), 7.62 (d, J=8.79 Hz, 1 H), 7.78 (m, 2 H), 8.18 (s, 1 H).
58	4-Iodo-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	394.2	395	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H) 7.64 (d, J=8.59 Hz, 2 H) 7.73 (dd, J=8.59, 1.95 Hz, 1 H) 7.80 (m, 1 H) 7.86 (d, J=8.59 Hz, 2 H) 7.94 (m, 1 H) 8.14 (d, J=1.95 Hz, 1 H)
59	4-Amino-N-(2-methyl-1,3-benzothiazol-5-yl)-3-nitro-benzamide	328.4	329	(400 MHz, DMSO-D6) δ ppm 2.79 (s, 3 H), 7.10 (d, J=8.98 Hz, 1 H), 7.75 (m, 1 H), 7.87 (s, 2 H), 7.96 (d, J=8.59 Hz, 1 H), 8.01 (dd, J=8.89, 2.25 Hz, 1 H), 8.39 (m, 1 H), 8.75 (d, J=2.15 Hz, 1 H), 10.37 (s, 1 H)
60	N-(2-Methyl-1,3-benzothiazol-5-yl)-4-vinyl-benzamide	294.4	295	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H) 5.40 (d, J=10.94 Hz, 1 H) 5.87 (d, J=17.57 Hz, 1 H) 6.78 (dd,

				J=17.57, 10.94 Hz, 1 H) 7.52 (d, J=8.20 Hz, 2 H) 7.78 (m, 2 H) 7.87 (d, J=8.40 Hz, 2 H) 7.99 (s, 1 H) 8.15 (s, 1 H)
61	4-Ethoxy-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	312.4	313	(400 MHz, chloroform-D) δ ppm 1.45 (t, J=7.03 Hz, 3 H) 2.83 (s, 3 H) 4.10 (q, J=14.06, 7.03 Hz, 2 H) 6.96 (d, J=8.98 Hz, 2 H) 7.77 (m, 2 H) 7.88 (d, J=6.83 Hz, 2 H) 7.98 (s, 1 H) 8.11 (m, 1 H)
62	4-Ethylsulfanyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	328.5	329	(400 MHz, chloroform-D) δ ppm 1.37 (t, J=7.42 Hz, 3 H) 2.83 (s, 3 H) 3.02 (q, J=14.65, 7.22 Hz, 2 H) 7.33 (d, J=8.79 Hz, 2 H) 7.76 (s, 2 H) 7.81 (d, J=8.59 Hz, 2 H) 8.13 (s, 2 H)
63	4-Dimethylamino-naphthalene-1-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide	361.5	362	(400 MHz, chloroform-D) δ ppm 2.85 (s, 3 H), 2.96 (m, 6 H), 7.03 (d, J=7.81 Hz, 1 H), 7.54 (m, 2 H), 7.72 (d, J=7.81 Hz, 1 H), 7.83 (m, 3 H), 8.13 (m, 1 H), 8.26 (m, 1 H), 8.43 (m, 1 H).

64	2-Fluoro-6-iodo-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	412.2	413	
65	4-Ethoxymethyl-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	326.4	327	(400 MHz, chloroform-D) δ ppm 1.27 (t, J=14.06, 7.03 Hz, 3 H) 2.84 (s, 3 H) 3.58 (q, J=14.06, 7.03 Hz, 2 H) 4.58 (s, 2 H) 7.47 (d, J=8.59 Hz, 2 H) 7.78 (m, 2 H) 7.88 (d, J=8.40 Hz, 2 H) 8.05 (s, 1 H) 8.14 (s, 1 H)
66	N-(2-Methyl-1,3-benzothiazol-5-yl)-4-trifluoromethoxy-benzamide	352.3	353	(400 MHz, chloroform-D) δ ppm 2.84 (s, 3 H) 7.33 (d, J=8.79 Hz, 2 H) 7.77 (m, 2 H) 7.95 (d, J=8.98 Hz, 2 H) 8.00 (s, 1 H) 8.14 (d, J=1.95 Hz, 1 H)
67	4-Chloro-3-fluoro-N-(2-methyl-1,3-benzothiazol-5-yl)-benzamide	320.8	321	(400 MHz, DMSO-D6) δ ppm 2.74 (s, 3 H), 7.70 (m, 1 H), 7.75 (m, 1 H), 7.82 (dd, J=8.40, 1.95 Hz, 1 H), 7.96 (m, 2 H), 8.396 (m, 1 H), 10.49 (s, 1 H)

68	4-tert-Butyl-N-(2-formyl-1,3-benzothiazol-5-yl)-benzamide	338.4	339	(400 MHz, DMSO-D6) δ 1.35 (s, 9 H), 7.27 (d, *J*=8.6 Hz, 2 H), 7.60-7.73 (m, 2H), 7.73 (d, *J*=8.6 Hz, 1 H), 7.86 (s, 1 H), 8.13 (s, 1 H), 8.84 (s, 1H)
69	4-tert-Butyl-N-(2-hydroxymethyl-1,3-benzothiazol-5-yl)-benzamide	340.5	341	(400 MHz, DMSO-D6) δ 1.30 (s, 9 H), 3.12 (s, 1H), 4.42 (s, 2H) 7.22 (d, *J*=8.6 Hz, 2 H), 7.62-7.76 (m, 2H), 7.83 (d, *J*=8.6 Hz, 1 H), 7.96 (s, 1 H), 8.25 (s, 1 H)

Example 70

4-tert-Butyl-N-(2-{[(2-methoxypyridin-3-yl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide.

A mixture of SeO_2 (4.44 g, 40.0 mmol) and 4-tert-butyl-N-(2-methyl-benzothiazol-5-yl)-benzamide (16.0 mmol) in dioxane (20.0 mL) was kept under a N_2 atmosphere and heated to 100 °C for 18 hours with vigorous stirring. After cooling to room temperature, the dioxane was removed by evaporation under reduced pressure. The resulting residue was dissolved in EtOAc, washed with brine, dried with Na_2SO_4, filtered and concentrated under reduced pressure to yield the aldehyde, MS (ESI$^+$) *m/z* 325.0 $[M+H]^+$. The aldehyde (100 mg, 0.300 mmol) was mixed with 2-methoxypyridin-3-amine (36.0 mg, 0.300 mmol) and $MgSO_4$ (100 mg) in THF (3.00 mL). After 18 hours, $B_{10}H_4$ (14.0 mg, 0.320 mmol) dissolved in MeOH (3.00 mL) was added. The mixture was stirred until the reaction appeared complete by TLC analysis. 1M NaOH was added and the solvents were evaporated. The residue was purified by flash chromatography eluting with mixtures of hexanes and EtOAc (4:1, 1:1). ^{1}H NMR (400 MHz, chloroform-D) δ ppm 1.29 (s, 9 H) 3.98 (s, 3 H) 4.68 (d, J=5.86 Hz, 2 H) 6.67 (m, 2 H) 7.40 (dt, J=8.69, 2.10 Hz, 2 H) 7.51

(dd, J=4.69, 1.95 Hz, 1 H) 7.66 (d, J=1.17 Hz, 2 H) 7.80 (ddd, J=8.59, 2.25, 2.05 Hz, 2 H) 8.25 (d, J=1.17 Hz, 1 H) 8.45 (s, 1 H); MS [M+H] calc. 447.2 found 447.0.

Example 71

4-tert-Butyl-N-[2-(1-hydroxyethyl)-1,3-benzothiazol-5-yl]benzamide

Methylmagnesium bromide (276 uL, 3.0 M in Et_2O) was added dropwise via syringe to a stirred solution of the aldehyde (obtained as an intermediate in Example 70) (100 mg, 0.30 mmol) in THF (10.0 mL) at -78°C under nitrogen. After addition was complete the mixture was stirred for additional 1 hour and quenched with saturated aqueous ammonium chloride (2.0 mL). The mixture was diluted with EtOAc (25.0 mL) and water (20.0 mL) and the organic phase was seaprated. The aqueous phase was extracted with EtOAc (2 X 10.0 mL) and the organic phases combined and washed with brine solution (30.0 mL). The organic was dried with Na_2SO_4, filtered and concentrated by rotary evaporator to a residue which was purified by column chromatography on silica gel using EtOAc/hexanes as an eluent to yield the title product. ^{1}H NMR (400 MHz, methanol-D4) δ ppm 1.34 (s, 9 H), 1.62 (d, J=6.44 Hz, 3 H), 5.12 (m, 1 H), 7.54 (d, J=8.59 Hz, 2 H), 7.69 (dd, J=8.69, 2.05 Hz, 1 H), 7.89 (m, 3 H), 8.39 (d, J=1.95 Hz, 1 H). MS [M+H] calc. 355.1 found 355.2.

Example 72

4-tert-Butyl-N-{2-[(1H-pyrazol-3-ylamino)methyl]-1,3-benzothiazol-5-yl}benzamide

The title compounds wa synthesized according to the procedure described in Example 70 using 1*H*-pyrazol-3-amine at the reductive amination step. ^{1}H NMR (400 MHz, methanol-D4) δ ppm 1.31 (m, 9 H) 4.71 (s, 2 H) 5.62 (d, J=2.34 Hz, 1 H) 7.35 (d, J=2.34 Hz, 1 H) 7.53 (d, J=8.79 Hz, 2 H) 7.67 (dd, J=8.69, 2.05 Hz, 1 H) 7.83 (d, J=8.59 Hz, 1 H) 7.88 (d, J=8.79 Hz, 2 H) 8.37 (d, J=1.76 Hz, 1 H); MS [M+H] calc. 406.2 found 406.0.

Example 73

4-(1,1-Dimethylethyl)-N-[2-[[(4-nitrophenyl)amino]methyl]-5-benzothiazolyl]-benzamide

The title compound was synthesized according to the procedure described in Example 70 using p-nitroaniline at the reductive amination step. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 1.32 (s, 9 H), 4.91 (m, 2 H), 6.78 (d, J=9.18 Hz, 2 H), 7.56 (d, J=8.40 Hz, 2 H), 7.76 (dd, J=8.79, 1.95 Hz, 1 H), 7.91 (d, J=8.40 Hz, 2 H), 7.97 (d, J=8.79 Hz, 1 H), 8.02 (d,

J=9.18 Hz, 2 H), 8.17 (t, J=6.25 Hz, 1 H), 8.50 (d, J=1.76 Hz, 1 H), 10.38 (s, 1 H). MS [M+H] calc. 461.2 found 461.0.

Example 74

N-[2-(Aminomethyl)-1,3-benzothiazol-5-yl]-4-tert-butylbenzamide

4-tert-Butyl-N-(2-hydroxymethyl-1,3-benzothiazol-5-yl)-benzamide (44.0 mg, 0.380 mmol) was mixed with MsCl (40.0 mg, 0.390 mmol, 0.0540 mL) and Et_3N (51.0 mg, 0.500 mmol) in DCM (5.00 mL) and the solution was stirred for 10 minutes. NH_3 (2.0M in EtOH) was added, and the mixture was stirred for additional 18 hours. The solvent was evaporated, and the crude product was purified by HPLC (Luna 15 u, C18 (2), 250 mm X 21.2 mm) eluting with mixtures of MeCN and H_2O containing 1%TFA. ^1H NMR (400 MHz, methanol-D4) δ ppm 1.21 (s, 9 H) 4.48 (s, 2 H) 7.41 (d, J=8.20 Hz, 2 H) 7.57 (d, J=8.20 Hz, 1 H) 7.76 (d, J=8.20 Hz, 2 H) 7.83 (d, J=8.59 Hz, 1 H) 8.47 (s, 1 H); MS [M+H] calc. 340.1 found 340.3.

Example 75

4-tert-Butyl-N-(2-{[(methylsulfonyl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide

N-[2-(Aminomethyl)-1,3-benzothiazol-5-yl]-4-*tert*-butylbenzamide (example 74) (130 mg, 0.384 mmol) was stirred with MsCl (44.0 mg, 0.387 mmol) and Et_3N (58.0 mg, 0.600 mmol, 0.0800 mL) in DCM (5.00 mL) for 1 hour. The solvent was evaporated, and the residue was purified by HPLC (Luna 15 u, C18 (2), 250 mm X 21.2 mm) eluting with mixtures of MeCN and H_2O containing 1%TFA to yield the title product. ^1H NMR (400 MHz, chloroform-D) δ ppm 1.35 (s, 9 H) 3.05 (s, 3 H) 4.79 (s, 2 H) 5.73 (s, 1 H) 7.52 (d, J=8.59 Hz, 2 H) 7.80 (s, 2 H) 7.85 (d, J=8.40 Hz, 2 H) 8.12 (s, 1 H) 8.23 (s, 1 H); MS [M+] calc. 417.5 found 417.9; Anal. found C 54.39% H 5.43% N 8.71%.

Example 76

N-{2-[(Acetylamino)methyl]-1,3-benzothiazol-5-yl}-4-tert-butylbenzamide

N-[2-(Aminomethyl)-1,3-benzothiazol-5-yl]-4-*tert*-butylbenzamide (example 74) (60.0 mg, 0.18 mmol) was stirred with acetyl chloride (16.0 mg, 0.2 mmol, 0.015 mL) and Et_3N (25.0 mg, 0.25 mmol) in DCM (5.00 mL) for 1 hour. The solvent was evaporated, and the residue was purified by HPLC eluting with mixtures of MeCN and H_2O containing 1%

TFA to yield the title product. ^{1}H NMR (400 MHz, chloroform-D) δ ppm 1.35 (s, 9 H) 2.12 (s, 3 H) 4.84 (s, 2 H) 7.49 (d, J=8.40 Hz, 2 H) 7.68 (s, 1 H) 7.75 (d, J=8.79 Hz, 1 H) 7.84 (d, J=8.20 Hz, 2 H) 8.20 (s, 1 H) 8.64 (s, 1 H) 11.35 (s, 1 H); MS [M+H] calc. 382.1 found 382.0; Anal. found C 55.85% H 4.94% N 8.60%.

Example 77

5-[(4-tert-Butylbenzoyl)amino]-1,3-benzothiazole-2-carboxamide

The aldehyde (example 70) (100 mg, 0.3 mmol) was dissolved in THF (10.0 mL) and a mixture of sodium chlorite (54.0 mg, 0.6 mmol) and sulfamic acid (58.0 mg, 0.6 mmol) in H_2O (5.0 mL) was added drop-wise. The mixture was stirred for 1 hour, and then the aqueous phase was extracted with EtOAc. The combined organic phases were dried over $MgSO_4$, filtered and evaporated to yield the acid, which was immediately dissolved in DCM (5.0 mL) containing a mixture of allyl chloroformate (48.0 mg, 0.400 mmol) and DMAP (48.0 mg, 0.400 mmol, 0.340 mL). The mixture was stirred for 1 hour and then evaporated to yield the mixed anhydride: MS [M+] calc. 435.0 found 435.9. The anhydride was dissolved in 5.0 mL of EtOH containing NH_3 (2.0M), and the mixture was stirred for 18 hours. The solvent was evaporated, and the product was purified by flash chromatography eluting with mixtures of hexanes and EtOAc (4:1, 1:1) to yield decarboxylated material (see example 77) and the title product. ^{1}H NMR (400 MHz, chloroform-D) δ ppm 1.34 (m, 9 H) 6.11 (s, 2 H) 7.41 (s, 1 H) 7.49 (d, J=7.62 Hz, 2 H) 7.73 (d, J=8.59 Hz, 1 H) 7.86 (d, J=7.23 Hz, 2 H) 8.33 (m, 1 H) 8.49 (s, 1 H); MS [M+H] calc. 354.1 found 354.0.

Example 78

N-1,3-Benzothiazol-5-yl-4-tert-butylbenzamide

See above (example 76). ^{1}H NMR (400 MHz, DMSO-D_6) δ ppm 1.30 (s, 9 H) 7.54 (d, J=8.40 Hz, 2 H) 7.82 (dd, J=8.79, 1.95 Hz, 1 H) 7.90 (d, J=8.59 Hz, 2 H) 8.08 (d, J=8.79 Hz, 1 H) 8.59 (d, J=1.95 Hz, 1 H) 9.36 (s, 1H) 10.38 (s, 1 H); IR (neat) 1661 cm^{-1}; MS [M+H] calc. 311.1 found 311.0.

Example 79

4-Chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide

According to amide bond forming procedure described in Example 2, 5-amino-2-methylbenzothiazole reacted with 4-chlorobenzoyl chloride to yield 4-chloro-N-(2-methyl-benzothiazol-5-yl)-benzamide: MS [M+] calc. 302, found 302.0. This intermediate was oxidized with SeO_2 to the corresponding aldehyde as described in Example 70. The aldehyde (3.30 mmol) was mixed with $NaBH_4$ (122 mg, 3.30 mmol) in MeOH (150 mL). After the reaction was complete according to TLC, the volatiles were removed and the residue was dissolved in a mixture of DCM and MeOH (10 mL, 1:5) and passed through a short pad of silica. The filtrate was concentrated and a residue was crystallized from a mixture of EtOAc and MeOH (40:1). A yellow solid formed was collected by filtration. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 4.85 (m, 2 H), 6.26 (t, J=5.96 Hz, 1 H), 7.62 (d, J=8.40 Hz, 2 H), 7.76 (m, 1 H), 8.02 (m, 3 H), 8.43 (m, 1 H), 10.50 (s, 1 H). MS [M+H] calc. 319.0 found 319.0.

Example 80

1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-propyl-1H-pyrazole-4-carboxamide

The title compound was synthesized from 5-amino-2-methylbenzothiazole and 1-(4-chlorophenyl)-5-propyl-1*H*-pyrazole-4-carbonyl chloride according to the procedure described in the example 79. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 0.76 (t, J=7.32 Hz, 3 H), 1.46 (m, 2 H), 2.97 (m, 2 H), 4.85 (d, J=6.05 Hz, 2 H), 6.26 (t, J=5.96 Hz, 1 H), 7.55 (d, J=8.79 Hz, 2 H), 7.65 (d, J=8.79 Hz, 2 H), 7.74 (dd, J=8.79, 1.95 Hz, 1 H), 8.01 (d, J=8.59 Hz, 1 H), 8.36 (m, 2 H), 10.07 (s, 1 H). MS [M+H] calc. 427.1 found 427.0.

Example 81

1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-(trifluoromethyl)-1H-pyrazole-4-carboxamide

The title compound was synthesized from 5-amino-2-methylbenzothiazole and 1-(4-chlorophenyl)-5-(trifluoromethyl)-1*H*-pyrazole-4-carbonyl chloride according to the procedure described in the example 79. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 4.86 (d, J=6.05 Hz, 2 H), 6.26 (t, J=5.96 Hz, 1 H), 7.60 (d, J=8.59 Hz, 2 H), 7.69 (m, 3 H), 8.04 (d,

J=8.59 Hz, 1 H), 8.37 (m, J=4.69 Hz, 2 H), 10.72 (s, 1 H). MS [M+H] calc. 453.0 found 452.9.

Example 82

N-(2,4-dimethyl-1,3-benzothiazol-5-yl)-4-(1-hydroxy-1-methylethyl)benzamide.

According to amide bond forming procedure described in Example 1, 5-amino-2-methylbenzothiazole reacted with 4-(methoxycarbonyl)benzoic acid to yield N-(2-Methyl-benzothiazol-5-yl)-terephthalamic acid methyl ester: MS [M+] calc. 326.0, found 326.0. This intermediate was placed into a flask, which was capped with a rubber septum and charged with N_2 gas. THF (10.0 mL) was added, followed by MeMgBr (4.60 mmol, 1.53 mL), and the reaction was stirred for 8 hours at room temperature. A saturated solution of NH_4Cl was added, and the mixture was evaporated to dryness in vacuum. The residue was purified by HPLC eluting with mixtures of MeCN and H_2O containing 1%TFA to yield the title product. ^1H NMR (400 MHz, CHLOROFORM-D) δ ppm 1.62 (s, 6 H) 2.96 (s, 3 H) 3.50 (s, 1 H) 7.62 (d, J=8.59 Hz, 2 H) 7.83 (d, J=8.79 Hz, 1 H) 7.90 (d, J=8.59 Hz, 2 H) 8.09 (dd, J=8.79, 1.76 Hz, 2 H) 8.21 (s, 1 H) 8.27 (s, 1 H); MMS [M+] cald. 327.1, found 327.0.

Example 83

4-(Hydroxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide

According to amide bond forming procedure described in Example 1, allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate reacted with *p*-carboxybenzaldehyde to yield carbonic acid allyl ester 5-(4-formyl-benzoylamino)-benzothiazol-2-ylmethyl ester. This intermediate (97 mg, 0.25 mmol) and $B_{10}H_{14}$ (30 mg, 0.25 mmol) were stirred in MeOH (10. 0 mL) for 48 hours. The reaction mixture was diluted with EtOAc (40.0 mL) and water (30.0 mL) and the organic phase separated. The aqueous phase was extracted with EtOAc (2 X 10.0 mL) and the combined organic phases were washed with brine solution (30.0 mL). The organic phase was dried with Na_2SO_4, filtered and concentrated in vacuum. The residue was purified by silica gel chromatography (1:1 EtOAc/hexanes) to yield the title product. ^1H NMR (400 MHz, DMSO-D6) δ ppm 4.56 (m, 2 H), 4.83 (m, 2 H), 5.32 (t, J=5.66 Hz, 1 H), 6.21 (t, J=5.96 Hz, 1 H), 7.45 (d, J=8.20 Hz, 2 H), 7.76 (dd,

J=8.69, 1.86 Hz, 1 H), 7.94 (d, J=8.20 Hz, 2 H), 7.99 (d, J=8.59 Hz, 1 H), 8.42 (d, J=1.76 Hz, 1 H), 10.35 (s, 1 H). MS [M+H] calc. 315.1 found 315.0.

Example 84

4-tert-butyl-N-(4-cyano-2-methyl-1,3-benzothiazol-5-yl)benzamide

N-(4-Bromo-2-methyl-1,3-benzothiazol-5-yl)-4-tert-butylbenzamide (example 12) (50.0 mg, 0.124 mmol) and CuCN (22 mg, 0.248 mmol) were dissolved in DMF (3.00 mL) and heated to 250 °C in a microwave oven for 20 minutes. The mixture was cooled, and the solvent was evaporated. The residue was purified by flash chromatography on silica gel eluting with mixtures of hexanes and EtOAc (4:1, 2:1, 1:1) to yield the title product. ^{1}H NMR (400 MHz, chloroform-D) δ ppm 1.36 (s, 9 H) 2.92 (s, 3 H) 7.55 (ddd, J=8.74, 2.25, 2.10 Hz, 2 H) 7.92 (ddd, J=8.64, 2.25, 2.00 Hz, 2 H) 8.03 (d, J=8.98 Hz, 1 H) 8.58 (s, 1 H) 8.69 (m, 1H); MS [M+] calcd. 350.1, found 350.0.

Example 85

4-tert-butyl-N-[2-(hydroxymethyl)-1,3-benzoxazol-5-yl]benzamide

A solution of 2-methyl-5-nitro-1,3-benzoxazole (500 mg, 2.8 mmol) in (dimethoxymethyl)dimethylamine (5.0 ml) was stirred in the microwave at 200°C for 15 min. (900 sec.). The precipitate was filtered off, washed with methanol and dried yielding (*E*)-*N*,*N*-dimethyl-2-(5-nitro-1,3-benzoxazol-2-yl)ethylenamine, 200 mg (31%), as a yellow powder. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 2.72 - 3.20 (m, 6 H) 5.02 (d, *J*=13.08 Hz, 1 H) 7.59 (d, *J*=8.79 Hz, 1 H) 7.74 (d, *J*=13.08 Hz, 1 H) 8.00 (dd, *J*=8.79, 2.34 Hz, 1 H) 8.10 (d, *J*=2.34 Hz, 1 H).

(*E*)-*N*,*N*-Dimethyl-2-(5-nitro-1,3-benzoxazol-2-yl)ethylenamine (200mg, 0.86 mmol) dissolved in methanol (20 ml), was hydrogenated over 10% palladium on carbon (500 mg) for 1 hour. The catalyst was removed via filtration through Celite and the filtrate was concentrated to yield a crude 2-[(*E*)-2-(dimethylamino)vinyl]-1,3-benzoxazol-5-amine, 120 mg (69%), which used as such in the next reaction step. ^{1}H NMR (400 MHz, DMSO-D6) δ ppm 2.88 (s, 6 H) 4.76 (s, 2 H) 4.88 (d, J=13.28 Hz, 1 H) 6.31 (dd, J=8.50, 2.25 Hz, 1 H) 6.54 (d, J=2.25 Hz, 1 H) 7.02 (d, J=8.50 Hz, 1 H) 7.48 (d, J=13.28 Hz, 1 H).

2-[(*E*)-2-(Dimethylamino)vinyl]-1,3-benzoxazol-5-amine (100 mg, 0.49 mmol) was dissolved in DCM (5.0 ml) containing dimethylaminopyridine (179 mg, 0.74 mmol). 4-

tert-Butylbenzoyl chloride (144mg, 1.47 mmol) was added and the mixture was stirred at ambient temperature for 1h. The mixture was diluted with DCM and extracted with water. The organic phase was dried over anhydrous sodium sulphate, and concentrated under reduced pressure. The residue was purified on a small silica gel column using ethyl acetate as the eluent to yield 4-*tert*-butyl-*N*-{2-[(*E*)-2-(dimethylamino)vinyl]-1,3-benzoxazol-5-yl}benzamide, 45 mg (25%). ^{1}H NMR (400 MHz, methanol-D4) δ ppm 1.37 (s, 9 H) 2.99 (s, 6 H) 5.02 (d, J=13.28 Hz, 1 H) 7.29 - 7.35 (m, 1 H) 7.35 - 7.44 (m, 1 H) 7.51 - 7.57 (m, 2 H) 7.63 (d, J=13.28 Hz, 1 H) 7.78 (d, J=1.95 Hz, 1 H) 7.83 - 7.93 (m, 2 H). 4-*tert*-Butyl-*N*-{2-[(*E*)-2-(dimethylamino)vinyl]-1,3-benzoxazol-5-yl}benzamide. (45mg, 0.124 mmol) was dissolved in a mixture of THF and water (1:1, 10 ml) and sodium periodate (158 mg, 0.74 mmol) was added. The mixture was stirred at ambient temperature for 3 h. The solution was extracted with DCM, the organic phase was dried over anhydrous sodium sulphate and concentrated under reduced pressure. The residue was dissolved in methanol (20 ml) and treated with sodium borohydride (200 mg, 5.4 mmol) at ambient temperature for 1 h. The reaction mixture was diluted with water and extracted with dichloromethane. The organic phase was dried over anhydrous sodium sulphate and concentrated under reduced pressure. The residue was purified by a column chromatography on silicagel using 50% ethyl acetate in hexane as an eluent to yield the title product, 18 mg (45%) as colourless oil. ^{1}H NMR (400 MHz, methanoL-D4) δ ppm 1.35 (s, 9 H) 4.79 (s, 2 H) 7.45 - 7.64 (m, 3 H) 7.61 - 7.73 (m, 1 H) 7.80 - 7.96 (m, 2 H) 8.12 (d, J=1.76 Hz, 1 H). MS [M+] calcd. 325.2, found 325.0.

Example 86

5-(4-tert-butylbenzoylamino)-1,3-benzothiazol-2-ylcarboxylic acid

A solution of 4-tert-butyl-*N*-(2-formyl-1,3-benzothiazol-5-yl)-benzamide (0.1 mmol) in THF (2 mL) was treated sequentially with a solution of sulfamic acid (0.2 mmol) in water (0.5 mL) and a solution of sodium chlorite (0.15 eq) in water (0.5 mL). The mixture was stirred at ambient temperature for 1 h, then partitioned between ethyl acetate (5 mL) and water (5 mL). The organic phase was separated, the water phase was extracted 3 times with ethyl acetate. Combined organic phase was dried over anhydrous sodium sulphate and concentrated *in vacuo* . The crude material was purified by preparative HPLC (X-Terra C8 column, 19x 300 mm), using a gradient of A (water 95%, containing NH_4OAc (0.01 M),

and 5% acetonitrile) and B (acetonitrile), to give the title compound. MS [M+] calcd. 354.4, found 355.0

Example 87

4-tert-Butyl-N-(2-methoxycarbonyl-1,3-benzothiazol-5-yl)-benzamide

A solution of 5-(4-tert-butylbenzoylamino)-1,3-benzothiazol-2-yl carboxylic acid (0.1 mmol) in methanol (3 mL) was treated with one drop of concentrated hydrochloric acid. The mixture was concentrated to dryness *in vacuo*. The oily residue was then purified by preparative HPLC (X-Terra C8 column, 19x 300 mm), using a gradient of A (water 95%, containing NH_4OAc (0.01 M), and 5% acetonitrile) and B (acetonitrile), to give the title compound as a solid. ^1H NMR (400 MHz, DMSO-d_6) δ 1.32 (s, 9 H), 3.65 (s, 1H), 7.25 (d, *J*=8.6 Hz, 2 H), 7.65-7.79 (m, 2H), 7.85 (d, *J*=8.6 Hz, 1 H), 7.91 (s, 1 H), 8.29 (s, 1 H). MS [M+] calcd. 368.5, found 369

Pharmacology

1. hVR1 FLIPR (Fluorometric Image Plate Reader) screening assay

Transfected CHO cells, stably expessing hVR1 (15,000 cells/well) are seeded in 50 ul media in a black clear bottom 384 plate (Greiner) and grown in a humidified incubator (37°C, 2% CO_2), 24-30 hours prior to experiment.

Subsequently, the media is removed from the cell plate by inversion and 2 μM Fluo-4 is added using a multidrop (Labsystems). Following the 40 minutes dye incubation in the dark at 37°C and 2% CO_2, the extracellular dye present is washed away using an EMBLA (Scatron), leaving the cells in 40ul of assay buffer (1 X HBSS, 10 mM D-Glucose, 1 mM $CaCl_2$, 10 mM HEPES, 10 X 7.5% $NaHCO_3$ and 2.5 mM Probenecid).

FLIPR assay - IC_{50} determination protocol

For IC_{50} determinations the fluorescence is read using FLIPR filter 1 (em 520-545 nM). A cellular baseline recording is taken for 30 seconds, followed by a 20 μl addition of 10, titrated half-log concentrations of the test compound, yielding cellular concentration ranging from 3 μM to 0.1 nM. Data is collected every 2 seconds for a further 5 minutes

prior to the addition of a VR1 agonist solution: either 50 nM solution of capsaicin or MES (2-[N-morpholino] ethanesulfonic acid) buffer (pH 5.2), by the FLIPR pipettor. The FLIPR continues to collect data for a further 4 minutes. Compounds having antagonistic properties against the hVR1 will inhibit the increase in intracellular calcium in response to the capsaicin addition. This consequently leading to a reduction in fluorescence signal and providing a reduced fluorescence reading, compared with no compound, buffer controls. Data is exported by the FLIPR program as a sum of fluorescence calculated under the curve upon the addition of capsaicin. Maximum inhibition, Hill slope and IC_{50} data for each compound are generated.

2. DRGs were dissected out from adult Sprague Dawley rats (100-300 gr), and placed on ice in L15 Leibovitz medium. The ganglia were enzyme treated with Collagenase 80U/ml+ Dispase 34 U/ml dissolved in DMEM +5% serum, overnight at 37 °C. The next day, cells were triturated with fire polished pasteur pipettes, and seeded in the center of 58 mm diameter Nunc cell dishes coated with Poly-D Lysine (1 mg/mL). The DRGs were cultured in a defined medium without foetal bovine serum, containing Dulbecco's MEM / NUT MIX F-12 (1:1) without L-glutamine but with pyridoxine, 6 mg/mL D(+)-Glucose, 100 µg/mL apo-transferrin, 1 mg/mL BSA, 20 µg/mL insulin, 2 mM L-glutamine, 50 IU/ mL Penicillin, 50 µg / mL Streptomycin and 0.01 µg/mL NGF-7S.

When the cells had grown for 2 days up to 4 weeks, the experiments were done. Cells were chosen based on size and presence of neurites. Small cells with long processes were used for recording (most likely to be C neurons, with native VR1 receptors).

The cells were recorded with conventional whole cell voltage clamp patch clamp, using the following solutions (calcium ion free):
The extracellular solution comprised (in mM): NaCl 137, KCl 5, $MgCl_2 * H_2O$ 1.2, HEPES 10, Glucose 10, EGTA 5, Sucrose 50, pH to 7.4 with NaOH.
The intracellular solution comprised K-gluconate 140, NaCl 3, $MgCl_2 * H_2O$ 1.2, HEPES 10, EGTA 1, pH to 7.2 with KOH. When the cells were penetrated with suction, a puff of capsaicin (500 nM) was used to determine if the cell expressed VR1 receptor. If not, a new

cell was chosen. If yes, then the compounds were added in increasing doses before the capsaicin pulse (500 nM), to determine an IC_{50} value.

List of abbreviations

VR1	vanilloid receptor 1
IBS	irritable bowel syndrome
IBD	inflammatory bowel disease
GERD	gastro-esophageal reflux disease
DRG	Dorsal Root Ganglion
BSA	Bovine Serum Albumin
HEPES	4-(2-Hydroxyethyl)piperazine-1-ethanesulfonic acid
EGTA	Ethylene glycol-bis(2-aminoethylether)-*N,N,N',N'*-tetraacetic acid
DMEM	Dulbeccos Modified Eagle's Medium

Results

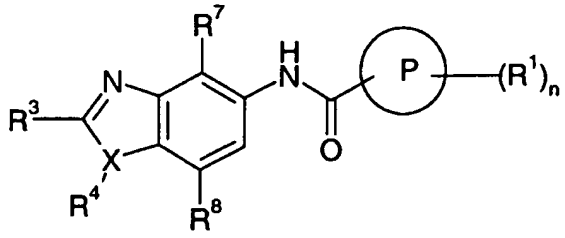
Typical IC_{50} values as measured in the assays described above are 10 μM or less. In one aspect of the invention the IC_{50} is below 500 nM. In another aspect of the invention the IC_{50} is below 100 nM. In a further aspect of the invention the IC_{50} is below 10 nM.

Results from the hVR1 FLIPR

Example No.	IC_{50} nM (agonist)	
2	10 (capsaicin)	60 (H^+/MES buffer)
71	200 (capsaicin)	
19	50 (capsaicin)	45 (H^+/MES buffer)

CLAIMS

1. A compound having the formula I



wherein:

ring P is C_{6-10}aryl, C_{3-7}cycloalkyl, C_{5-6}heteroaryl, which ring P may be fused with phenyl, C_{5-6}heteroaryl, C_{3-7}cycloalkyl or C_{3-7}heterocycloalkyl;

R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, C_{1-6}alkyl, C_{2-6}alkenyl, C_{2-6}alkynyl, C_{1-6}haloalkyl, C_{1-6}haloalkylO, phenylC_{0-6}alkyl, C_{5-6}heteroarylC_{0-6}alkyl, C_{3-7}cycloalkylC_{0-6}alkyl, C_{3-7}heterocycloalkylC_{0-6}alkyl, C_{1-6}alkylOC_{0-6}alkyl, C_{1-6}alkylSC_{0-6}alkyl or C_{1-6}alkylNC_{0-6}alkyl;

n is 1, 2, 3, 4 or 5;

X is O or S, when

R^3 is H, C_{1-6}alkyl, C_{1-6}haloalkyl, R^5OC_{1-6}alkyl, R^5OCO, R^5CO, NR^5R^6CO, $NR^5R^6C_{0-6}$alkyl, C_{2-6}alkenylOC_{0-6}alkyl or hydroxyC_{1-6}alkyl; and

R^4 is nil; or

X is N, when

R^3 is H, C_{1-6}alkyl, C_{1-6}iodoalkyl, C_{1-6}bromoalkyl, C_{1-6}chloroalkyl, C_{1-6}alkylOC_{0-6}alkyl, R^5OC_{1-6}alkyl, R^5CO, R^5CO2, NR^5R^6CO, $NR^5R^6C_{0-6}$alkyl or C_{2-6}alkenylOC_{0-6}alkyl; and

R^4 is H, C_{1-4}alkyl, hydroxyC_{1-6}alkyl or C_{1-6}alkylOC_{1-6}alkyl; or

X is N, when R^3 is C_{1-6}fluoroalkyl or hydroxyC_{1-2}alkyl and R^4 is H;

R^5 and R^6 are independently selected from H, C_{1-6}alkyl, C_{6-10}aryl, C_{5-6}heteroaryl, C_{1-4}alkylSO_2 and C_{1-3} alkylCO;

R^7 and R^8 are independently selected from H, C_{1-6}alkyl, halo, cyano, C_{1-6}alkylOC_{0-6}alkyl, OH, NO_2 and COR^9, $N(R^9)_2$;

R^9 is H or C_{1-6}alkyl;

and wherein any alkyl, alkylOalkyl, haloalkyl, haloalkylO, phenyl, heteroaryl, cycloalkyl or heterocycloalkyl group may be substituted with one or more A; and

A is OH, NO_2, R^9CO, $R^9O(CO)$, $N(R^9)_2$, R^9S, R^9SO_2, halo or C_{1-6}alkylOC_{0-6}alkyl,
or salts, solvates or solvated salts thereof.

2. The compound according to claim 1 wherein
ring P is C_{6-10}aryl, C_{5-6}heteroaryl, which ring P may be fused with
C_{3-7}heterocycloalkyl;
R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, C_{1-6}alkyl, C_{2-6}alkenyl, C_{1-6}haloalkyl,
C_{1-6}haloalkylO, phenylC_{0-6}alkyl, C_{3-7}heterocycloalkylC_{0-6}alkyl, C_{1-6}alkylOC_{0-6}alkyl or
C_{1-6}alkylSC_{0-6}alkyl;
n is 1, 2 or 3;
X is O or S, when
R^3 is C_{1-6}alkyl, NR^5R^6CO, $NR^5R^6C_{0-6}$alkyl, C_{2-6}alkenylOC_{0-6}alkyl or hydroxyC_{1-6}alkyl; and
R^4 is nil; or
X is N, when
R^3 is H or C_{1-6}alkyl; and
R^4 is C_{1-4}alkyl or hydroxyC_{1-6}alkyl; or
X is N, when R^3 is C_{1-6}fluoroalkyl and R^4 is H;
R^5 and R^6 are independently selected from H, C_{6-10}aryl, C_{5-6}heteroaryl,
C_{1-4}alkylSO_2 and C_{1-3} alkylCO;
R^7 and R^8 are independently selected from H, halo and cyano;
and wherein any alkyl, phenyl, heteroaryl group may be substituted with one or more A;
and
A is OH, NO_2, halo or C_{1-6}alkylOC_{0-6}alkyl.

3. The compound according to any one of claims 1 or 2 wherein X is S and R^3 is methyl.

4. The compound according to any one of claims 1 or 2 wherein X is O and R^3 is C_{1-6}alkyl
or hydroxyC_{1-6}alkyl.

5. The compound according to any one of claims 1 or 2 wherein X is N and R^3 is C_{1-6}alkyl
and R^4 is C_{1-6}alkyl or hydroxyC_{1-6}alkyl.

6. The compound according to any one of claims 1 to 5 wherein
ring P is phenyl and R^1 is NO_2, NH_2, halo, $N(C_{1-6}alkyl)_2$, $C_{1-6}alkyl$, $C_{2-6}alkenyl$, $C_{2-6}alkynyl$, $C_{1-6}haloalkyl$, $C_{1-6}haloalkylO$, $phenylC_{0-6}alkyl$, $C_{5-6}heteroarylC_{0-6}alkyl$, $C_{3-7}cycloalkylC_{0-6}alkyl$, $C_{3-7}heterocycloalkylC_{0-6}alkyl$, $C_{1-6}alkylOC_{0-6}alkyl$, $C_{1-6}alkylSC_{0-6}alkyl$ or $C_{1-6}alkylNC_{0-6}alkyl$ optionally substituted with one or more A.

7. The compound according to any one of claims 1 to 5 wherein
ring P is pyrazolyl, pyridine, benzdioxolane, furan, thiophene or naphthalene.

8. Compounds selected from the group consisting of
3-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,
2-tert-Butyl-5-methyl-2*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
2-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,
2-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide,
4-Fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-trifluoromethyl-benzamide,
3,4-Dimethyl-*N*-(2-methyl-benzothiazol-5-yl)-benzamide,
2,2-Difluoro-benzo[1,3]dioxole-5-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
N-(2-Methyl-1,3-benzothiazol-5-yl)-6-trifluoromethyl-nicotinamide,
N-(2-Methyl-1,3-benzothiazol-5-yl)-4-propyl-benzamide,
3-Iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,
2,5-Dimethyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
5-tert-Butyl-2-methyl-furan-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
4-Bromo-3-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,
3,4-Difluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,
3-Chloro-2-fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,
Pyridine-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
2-Benzyl-5-tert-butyl-2*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,
3-Fluoro-4-trifluoromethyl-*N*-(2-trifluoromethyl-1*H*-benzimidazol-5-yl)-benzamide,
2-Fluoro-5-trifluoromethyl-*N*-(2-trifluoromethyl-1*H*-benzimidazol-5-yl)-benzamide,
4-Chloro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

1-Phenyl-5-trifluoromethyl-1*H*-pyrazole-3-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

1-Phenyl-5-propyl-1*H*-pyrazole-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2,3-Difluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-4-trifluoromethyl-benzamide,

3-Fluoro-4-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-methyl-benzooxazol-5-yl)-benzamide,

Biphenyl-4-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

3-Bromo-thiophene-2-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

4-Bromo-2-methyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

2-Chloro-3,4-dimethoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Amino-*N*-(2-methyl-1,3-benzothiazol-5-yl)-3-nitro-benzamide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-4-vinyl-benzamide,

4-Ethoxy-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethylsulfanyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Dimethylamino-naphthalene-1-carboxylic acid (2-methyl-1,3-benzothiazol-5-yl)-amide,

2-Fluoro-6-iodo-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-Ethoxymethyl-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

N-(2-Methyl-1,3-benzothiazol-5-yl)-4-trifluoromethoxy-benzamide, and

4-Chloro-3-fluoro-*N*-(2-methyl-1,3-benzothiazol-5-yl)-benzamide,

4-*tert*-Butyl-*N*-(2-formyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butyl-*N*-(2-hydroxymethyl-1,3-benzothiazol-5-yl)-benzamide,

5-(4-tert-butylbenzoylamino)-1,3-benzothiazol-2-ylcarboxylic acid, and

4-tert-Butyl-*N*-(2-methoxycarbonyl-1,3-benzothiazol-5-yl)-benzamide,

4-tert-Butoxy-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

4-Bromo-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-iodobenzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-morpholin-4-ylbenzamide,

N-{2-[(Allyloxy)methyl]-1,3-benzothiazol-5-yl}-4-morpholin-4-ylbenzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-propyl-1H-pyrazole-4-carboxamide,

1-tert-Butyl-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-3-methyl-1H-pyrazole-5-carboxamide,

4-(Ethoxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-1H-pyrazole-5-carboxamide,

4-Bromo-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-2-methylbenzamide,

4-tert-Butoxy-N-(2-methyl-1,3-benzoxazol-5-yl) benzamide,

N-(4-Bromo-2-methyl-1,3-benzothiazol-5-yl)-4-tert-butylbenzamide,

4-tert-Butyl-N-(4,7-dibromo-2-methyl-1,3-benzothiazol-5-yl)benzamide,

N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-1-phenyl-5-(trifluoromethyl)-1H-pyrazole-4-carboxamide,

4-Iodo-N-(2-methyl-5-benzothiazolyl)benzamide,

4-(tert-Butoxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

N-(1,2-Dimethyl-1H-benzimidazol-5-yl)-4-iodobenzamide,

4-tert-Butyl-N-(2-{[(2-methoxypyridin-3-yl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide,

4-tert-Butyl-N-[2-(1-hydroxyethyl)-1,3-benzothiazol-5-yl]benzamide,

4-tert-Butyl-N-{2-[(1H-pyrazol-3-ylamino)methyl]-1,3-benzothiazol-5-yl}benzamide,

4-(1,1-Dimethylethyl)-N-[2-[[(4-nitrophenyl)amino]methyl]-5-benzothiazolyl]-benzamide,

N-[2-(Aminomethyl)-1,3-benzothiazol-5-yl]-4-tert-butylbenzamide,

4-tert-Butyl-N-(2-{[(methylsulfonyl)amino]methyl}-1,3-benzothiazol-5-yl)benzamide,

N-{2-[(Acetylamino)methyl]-1,3-benzothiazol-5-yl}-4-tert-butylbenzamide,

5-[(4-tert-Butylbenzoyl)amino]-1,3-benzothiazole-2-carboxamide,

N-1,3-Benzothiazol-5-yl-4-tert-butylbenzamide,

4-Chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,

1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-propyl-1H-pyrazole-4-carboxamide,

1-(4-chlorophenyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-5-(trifluoromethyl)-1H-pyrazole-4-carboxamide,

N-(2,4-dimethyl-1,3-benzothiazol-5-yl)-4-(1-hydroxy-1-methylethyl)benzamide,

4-(Hydroxymethyl)-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide and

4-tert-butyl-N-(4-cyano-2-methyl-1,3-benzothiazol-5-yl)benzamide,
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-[2,2,2-trifluoro-1-hydroxy-1-(trifluoromethyl)ethyl]benzamide,
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-isopropoxybenzamide,
4-Bromo-2-chloro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
4-Bromo-2-fluoro-N-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
N-[2-(Hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(morpholin-4-ylmethyl)benzamide,
3-Fluoro-*N*-[2-(hydroxymethyl)-1,3-benzothiazol-5-yl]-4-(trifluoromethyl)benzamide,
4-tert-butoxy-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
4-(tert-Butoxymethyl)-N-[4-chloro-2-(hydroxymethyl)-1,3-benzothiazol-5-yl]benzamide,
and
4-*tert*-butyl-*N*-[2-(hydroxymethyl)-1,3-benzoxazol-5-yl]benzamide,
or salts, solvates or solvated salts thereof.

9. A pharmaceutical composition comprising as active ingredient a therapeutically effective amount of the compound according to any one of claims 1 to 8, in association with one or more pharmaceutically acceptable diluents, excipients and/or inert carriers.

10. The pharmaceutical composition according to claim 9, for use in the treatment of VR1 mediated disorders and for treatment of acute and chronic pain disorders, acute and chronic neuropathic pain and acute and chronic inflammatory pain.

11. The compound according to any one of claims 1 to 8, for use in therapy.

12. Use of the compounds of formula I according to any one of claims 1 to 8, in the manufacture of a medicament for treatment of VR1 mediated disorders.

13. The use according to claim 12 for treatment of acute and chronic pain disorders.

14. The use according to claim 12 for treatment of acute and chronic neuropathic pain.

15. The use according to claim 12 for treatment of acute and chronic inflammatory pain.

16. The use according to claim 12 for treatment of arthritis, fibromyalgia, low back pain, post-operative pain, visceral pains like chronic pelvic pain, cystitis, irritable bowel syndrome (IBS), pancreatitis, ischeamic, sciatia, diabetic neuropathy, multiple sclerosis, interstitial cystitis and pain related to interstitial cystitis, HIV neuropathy, asthma, cough and inflammatory bowel disease (IBD), gastro-esophageal reflux disease (GERD), psoriasis, cancer, emesis, urinary incontinence and hyperactive bladder.

17. The use according to claim 12 for treatment of respiratory diseases.

18. A method of treatment of VR1 mediated disorders and for treatment of acute and chronic pain disorders, acute and chronic neuropathic pain and acute and chronic inflammatory pain, and respiratory diseases, comprising administrering to a mammal, including man in need of such treatment, a therapeutically effective amount of the compounds of formula I, according to any one of claims 1 to 8.

19. Compounds selected from the group consisiting of
allyl (5-amino-1,3-benzothiazol-2-yl)methyl carbonate,
4-tert-Butyl-N-(2-formyl-1,3-benzothiazol-5-yl)-benzamide, and
4-Bromo-2-methyl-benzothiazol-5-ylamine, and
4-chloro-2-methyl-benzothiazole-5-ylamine.

20. Use of compounds according to claim 19 as intermediates in the preparation of the compound of formula I.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 2004/000635

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 277/64, C07D 417/12, C07D 235/08, C07D 235/10, C07D 277/68, C07D 263/56, A61K 31/4184, A61K 31/5377, A61P 25/04, A61P 25/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K, A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CHEM.ABS.DATA, WPI DATA, EPO-INTERNAL

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
E,X	WO 2004056774 A2 (NEUROGEN CORPORATION), 8 July 2004 (08.07.2004) --	1-18
P,X	WO 03037274 A2 (ICAGEN, INC.), 8 May 2003 (08.05.2003) --	1,2,5-7, 9-16,18
P,X	WO 2004011440 A1 (BANYU PHARMACEUTICAL CO., LTD), 5 February 2004 (05.02.2004) --	1,6

[X] Further documents are listed in the continuation of Box C. [X] See patent family annex.

* Special categories of cited documents:
"A" document defining the general state of the art which is not considered to be of particular relevance
"E" earlier application or patent but published on or after the international filing date
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"O" document referring to an oral disclosure, use, exhibition or other means
"P" document published prior to the international filing date but later than the priority date claimed
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
20 Sept 2004	21 -09- 2004
Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Facsimile No. +46 8 666 02 86	Authorized officer ANNA SJÖLUND/BS Telephone No. +46 8 782 25 00

Form PCT/ISA/210 (second sheet) (January 2004)

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, File REGISTRY, see RN: 627074-44-6, 626202-75-3, 540516-48-1, 503429-34-3, 498535-73-2, 497247-81-1, 497242-15-6, 489464-52-0, 477554-52-2, 477506-36-8, 392251-13-7, 391223-69-1, 372081-90-8, 338431-35-9, 335397-26-7, 335397-24-5, 330567-86-7, 330189-20-3, 314027-99-1, 313500-75-3, 300572-54-7, 300572-47-8, 497240-67-2, 477554-53-3, 477492-97-0, 476284-34-1, 391218-49-8, 381696-06-6, 335397-25-6, 330189-19-0, 313959-23-8, 293766-14-0, January 2001 --	1-3,5-7
X	STN International, file ACS, ACS accession no. 132:265201, Suzuki, Nobuhiro; Takekawa, Shiro; Kubo, Keiji; Imaeda, Yasuhiro (Takeda Chemical Industries, Ltd., "Preparation of imidazole derivatives as gonadotropinreleasing hormone antagonists", & JP,A2,2000095767,20000404, 79 pp. CAS RN: 263022-44-2 --	1-12,16
X	STN International, file ACS, ACS accession no. 1999:626041, document no. 131:257447, Shinkai, Hisashi; Ito, Takao; Yamada, Hideki (Japan Tobacco Inc., Japan), "Preparation of amide derivatives as nociception antagonists", & WO,A1,9948492,19990930, 113 pp. CAS RN: 244219-85-0 --	1-12,16
X	WO 03014064 A1 (BAYER AKTIENGESELLSCHAFT), 20 February 2003 (20.02.2003) --	1-17
X	US 3711608 A (CAMPBELL), 16 January 1973 (16.01.1973) --	1-16
X	WO 0242273 A2 (BRISTOL-MYERS SQUIBB COMPANY), 30 May 2002 (30.05.2002), ex. 175,178,181 --	1-17

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 2004/000635

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, file CAPLUS, CAPLUS accession no. 1986:472101, document no. 105:72101, Miller, Robert E.; Reid Willis, A., Jr.(Dep. Parasitol., Walter Reed Army Inst. Res., Washington, DC, 20307, USA), "Schistosoma mansoni: salicylanilides as topical prophylactic against cercarial penetration of mice" & Experimental Parasitology, 61(3), 359-68, (English) 1956, CAS RN: 98290-34-7 ---	1,6,9-11
X	STN International, file CAPLUS, CAPLUS accession no. 1981:444733, document no. 95:444733, Frey, Christoph (Ciba-Geigy A.G., Switz). "Iminoisoindolinone metal complexes, method for pigmenting an organic high-molecular-weight material and high-molecular-weight organic material containing such a metal complex", & EP,23889,19810211 --- --------	1,6

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE 2004/000635

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 18
because they relate to subject matter not required to be searched by this Authority, namely:

see extra sheet

2. ☒ Claims Nos.: 1-2,4-5,7
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

see extra sheet

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see extra sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-17

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

Box II.1

Claim 18 relates to methods of treatment of the human or animal body by surgery or by therapy or diagnostic methods practiced on the human or animal body (Rule 39.1(iv)). Nevertheless, a search has been executed for this claim. The search has been based on the alleged effects of the compounds or compositions

Box II.2

The initial phase of the search revealed a very large number of documents relevant to the issue of novelty. So many documents were retrieved that it is impossible to determine which parts of the claims 1-2, 4-5, 7 may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons, a meaningful search over the whole breadth of the claims is impossible. Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to the compounds mentioned in claim 8, with due regard to the general idea underlying the present invention.

Present claim 12 relates to the treatment of diseases which are actually not well defined. The use of the definition "VR1 mediated disorders" in the present context is considered to lead to a lack of clarity within the meaning of Article 6 PCT. The lack of clarity is such as to render a meaningful search not fully possible.
Consequently, the search has been restricted to the parts relating to the diseases mentioned in claims in claims 13-18 with due regard to the general idea underlying the present invention.

Box III

The International Search Authority considers that there are 2 inventions covered by the claims indicated as follows:

I: Claims: 1-17, directed to compounds of formula I, compositions and uses thereof.

II: Claims 19-20, directed to intermediate compounds, and uses thereof in the preparation of the compound of formula I.

WO	2004056774	A2	08/07/2004	NONE			
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